



OIML TC 3/SC 5
February 2006

First Committee Draft - 1CD

Guide
for the application of ISO/IEC 17025
to assessment of Testing Laboratories
involved in legal metrology testing

EXPLANATORY NOTE

The first Working Draft (IWD) had been drawn up on the basis of the document numbered OIML MAA 02 accepted by the R 60 and R 76 Committees on Participation Review (CPR) and used for conducting the peer assessments decided on by the CPR.

This first Committee Draft (ICD) has been drawn up on the basis of the first Working Draft (IWD) and of the comments received from TC 3/SC5.

In addition the ICD includes some amendments which take into account the necessary clarification of the terminology which will be introduced in the revision of the OIML B 3 “OIML Certificate System for measuring Instruments” and OIML B 10-1 “Framework for a Mutual Acceptance Arrangement on OIML Type Evaluations”.

This document aims to be a guide for assessing the conformity to ISO/IEC 17025 of any Testing Laboratories involved in legal metrology testing. These Testing laboratories may be:

- subcontracting Testing laboratories of a national type approval body;*
- subcontracting Testing Laboratories of an inspection body, for instance responsible for initial verification;*
- subcontracting Testing Laboratories of an OIML Issuing Authority when implementing the OIML Mutual Acceptance Arrangement (MAA).*

This document is also intended to be used by peer assessment or accreditation auditors when interpreting ISO/IEC 17025 for legal metrology applications in connection with the OIML Mutual Acceptance Arrangement (MAA).

This document is being developed in cooperation with ILAC, which is a liaison, and the purpose is to have this guide adopted by both organizations (OIML and ILAC). Consequently, the same document could be used for accreditation under the scope of legal metrology and for peer assessment.

FOREWORD

The International Organization of Legal Metrology (OIML) is a worldwide, intergovernmental organization whose primary aim is to harmonize the regulations and metrological controls applied by the national metrological services, or related organizations, of its Member States. The two main categories of OIML publications are:

International Recommendations (OIML R), which are model regulations that establish the metrological characteristics required of certain measuring instruments and which specify methods and equipment for checking their conformity; the OIML Member States shall implement these Recommendations to the greatest possible extent;

International Documents (OIML D), which are informative in nature and intended to improve the work of the metrological services;

International Basic Publications (OIML B), which define the operating rules of the various OIML structures and systems;

International Guides (OIML G), which are informative in nature and which are intended to give guidelines for the application of certain requirements to legal metrology.

OIML Draft Recommendations, Documents and Guides are developed by Technical Committees or Subcommittees which are formed by the Member States. Certain international and regional institutions also participate on a consultation basis. Cooperative agreements are established between OIML and certain institutions, such as ISO and IEC, with the objective of avoiding contradictory requirements; consequently, manufacturers and users of measuring instruments, test laboratories, etc. may apply simultaneously OIML publications and those of other institutions.

International Recommendations, International Documents and International Guides are published in French (F) and English (E) and are subject to periodic revision.

This publication - reference OIML G XX, edition 200X (E) - was developed by the OIML Technical TC 3/SC 5 *Conformity assessment*. It was approved for final publication by the International Committee of Legal Metrology in 200X.

OIML Publications may be downloaded from the OIML web site in the form of PDF files. Additional information on OIML Publications may be obtained from the Organization's headquarters:

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| <p style="text-align: center;">ISO 17025 General requirements for the competence of testing and calibration laboratories to testing laboratory activities</p> | <p style="text-align: center;">Specific requirements and comments related to Testing Laboratories involved in legal metrology testing</p> |
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| <p>Introduction</p> | |
| <p>The first edition (1999) of this International Standard was produced as the result of extensive experience in the implementation of ISO/IEC Guide 25 and EN 45001, both of which it replaced. It contained all of the requirements that testing and calibration laboratories have to meet if they wish to demonstrate that they operate a management system, are technically competent, and are able to generate technically valid results.</p> <p>The first edition referred to ISO 9001:1994 and ISO 9002:1994. These standards have been superseded by ISO 9001:2000, which made an alignment of ISO/IEC 17025 necessary. In the second edition, clauses have been amended or added only when considered necessary in the light of ISO 9001:2000.</p> <p>Accreditation bodies that recognize the competence of testing and calibration laboratories should use this International Standard as the basis for their accreditation. Clause 4 specifies the requirements for sound management. Clause 5 specifies the requirements for technical competence for the type of tests and/or calibrations the laboratory undertakes.</p> <p>Growth in use of management systems generally has increased the need to ensure that laboratories which form part of larger organisations or offer other services can operate to a quality management system that is seen as compliant with ISO 9001 as well as with this Standard. Care has been taken, therefore, to incorporate all those requirements of ISO 9001 that are relevant to the scope of testing and calibration services that are covered by the laboratory's management system.</p> <p>Testing and calibration laboratories that comply with this International Standard will therefore also operate in accordance with ISO 9001.</p> <p>Conformity of the quality management system within which the laboratory operates to the requirements of ISO 9001 does not of itself demonstrate the competence of the laboratory to produce technically valid data and results. Nor does demonstrated conformity to this International Standard imply conformity of the quality management system within which the laboratory operates to all the requirements of ISO 9001.</p> <p>The acceptance of testing and calibration results between countries should be facilitated if laboratories comply with this International Standard and if they obtain accreditation from bodies which have entered into mutual recognition agreements with equivalent bodies in other countries using this International Standard.</p> | <p>This Guide complements the requirements of ISO/IEC 17025 in order to provide confidence in the laboratory which performs the required tests and examinations.</p> <p>For the purpose of this Guide, the relevant terms and definitions given in chapter 3 apply.</p> <p>For the implementation of the MAA, the tests and examinations to be considered are those required by the relevant OIML Recommendation and, if applicable, the additional tests validated by the CPR. The acceptance of testing results between countries is formalized through the Declarations of Mutual Confidence (DoMCs).</p> |

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| <p>The use of this International Standard will facilitate cooperation between laboratories and other bodies, and assist in the exchange of information and experience, and in the harmonisation of standards and procedures.</p> | |
| <p>1 Scope</p> | |
| <p>1.1 This International Standard specifies the general requirements for the competence to carry out tests and/or calibrations including sampling.</p> <p>It covers testing and calibration performed using standard methods, non-standard methods, and laboratory-developed methods.</p> | <p>The requirements of this Guide are applicable to all Testing Laboratories involved in legal metrology and in particular those involved in type evaluation tests and examinations.</p> <p>For the OIML Certificate System and for the implementation of the MAA, these Testing Laboratories are designated by the Issuing Authority (see 3.3.1 of OIML B 3 and 3.13 of OIML B 10-1).</p> <p>The Issuing Authority is responsible for selecting the sample(s) to be tested and, when applicable, for defining the tests and examinations to be carried out.</p> <p>Testing methods covered are those defined in the relevant OIML Recommendation and in the DoMC for additional tests validated by the CPR.</p> |
| <p>1.2 This International Standard is applicable to all organizations performing tests and/or calibration activities. These include, for example, first-, second- and third-party laboratories, and laboratories where testing and/or calibration forms part of inspection and product certification.</p> <p>This International Standard is applicable to all laboratories regardless of the number of personnel or the extent of the scope of testing and/or calibration activities. When a laboratory does not undertake one or more of the activities covered by this International Standard, such as sampling and the design/development of new methods, the requirements of those clauses do not apply.</p> | <p>In case of the implementation of the MAA, the requirements of this Guide are applicable to any body that performs tests and examinations under the DoMC (e.g. Testing Laboratories) and Issuing Authorities when they carry out tests and/or examinations.</p> |
| <p>1.3 The notes given provide clarification of the text, examples and guidance. They do not contain requirements and do not form an integral part of this International Standard.</p> | |
| <p>1.4 This International Standard is for use by laboratories in developing their management system for quality, administrative and technical operations. Laboratory customers, regulatory authorities and accreditation bodies may also use it in confirming or recognizing the competence of laboratories.</p> <p><i>NOTE 1: The term "management system" in this International Standard means the quality, administrative and technical systems that govern the operations of a laboratory.</i></p> | |

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| <p><i>NOTE 2: Certification of a management system is sometimes also called registration.</i></p> | |
| <p>1.5 Compliance with regulatory and safety requirements on the operation of laboratories is not covered by this International Standards.</p> | |
| <p>1.6 If testing and calibration laboratories comply with the requirements of this International Standard, they will operate a quality management system for their testing and calibration activities that also meets the principles of ISO 9001. Annex A provides nominal cross-references between this International Standard and ISO 9001. This International Standard covers technical competence requirements that are not covered by ISO 9001.</p> <p><i>NOTE 1:</i> <i>It might be necessary to explain or interpret certain requirements in this International Standard to ensure that the requirements are applied in a consistent manner. Guidance for establishing applications for specific fields, especially for accreditation bodies (see ISO/IEC 17011) is given in annex B.</i></p> <p><i>NOTE 2:</i> <i>If a laboratory wishes accreditation for part or all of its testing and calibration activities, it should select an accreditation body that operates in accordance with ISO/IEC 17011</i></p> | |
| <p>2 Normative references</p> | |
| <p>The following referenced documents are indispensable for the application of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.</p> <p>ISO/IEC 17000, Conformity assessment - Vocabulary and general principles</p> <p>VIM, <i>International vocabulary of basic and general terms in metrology, issued by BIPM, IEC, IFCC, ISO, IUPAC, IUPAP and OIML.</i></p> <p><i>NOTE: Further related standards, guides, etc. on subjects included in this International Standard are given in the bibliography.</i></p> | <p>In addition:</p> <p>OIML V 2 International vocabulary of basic and general terms in metrology (VIM) (bilingual French-English), 1993</p> <p>OIML G 1: Guide to the expression of uncertainty in measurement (GUM), 1997</p> <p>OIML B 3: OIML Certificate System for Measuring Instruments, 2003</p> <p>OIML B 10-1: Framework for a Mutual Acceptance Arrangement on OIML type evaluations, 2004</p> <p>OIML B 10-2: Checklists for Issuing Authorities and Testing Laboratories carrying out OIML Type Evaluations, 2004</p> <p>OIML D 19: Pattern evaluation and pattern approval, 1988</p> <p>OIML D 14: Training and qualification of legal metrology personnel, 2004</p> |

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| <p>3 Terms and Definitions</p> | |
| <p>3.1 For the purpose of this document, the relevant terms and definitions given in ISO/IEC 17000 and VIM apply.</p> <p><i>NOTE :</i> <i>General definitions related to quality are given in ISO 9000, whereas ISO/IEC 17000 gives definitions specifically related to, certification and laboratory accreditation. Where different definitions are given in ISO 9000, the definitions in ISO/IEC 17000 and VIM are preferred.</i></p> | <p>Type (pattern) evaluation (VIML 2.5) Systematic examination and testing of the performance of one or more specimens of an identified type (pattern) of measuring instruments against documented requirements, the results of which are contained in the evaluation report, in order to determine whether the type may be approved.</p> <p><i>NOTE</i> “Pattern” is used in legal metrology with the same meaning as “type”; in the entries below, only “type” is used.</p> <p>Examination (OIML B 3) Visual inspection of an instrument or device and relevant documentation to ensure that some specified requirements are met.</p> <p>Conformity (OIML B 3) Fulfilment by a measuring instrument type of metrological and technical requirements as specified in the relevant Recommendation.</p> <p>Issuing Authority Any Authority which is responsible for issuing Certificates within a specified system of legal metrology control. <i>Note: In case of type evaluation, an Issuing Authority may be a national Type Approval Body and/or, for the implementation of the OIML Mutual Acceptance Arrangement (MAA), an OIML Issuing Authority according to 2.13 of OIML B 3 and 3.11 of B 10-1.</i></p> <p>Testing Laboratory Any necessary specialized laboratory or laboratories designated by the Issuing Authority to carry out examination and testing of a sample or samples of a measuring instrument submitted for evaluation, with the Issuing Authority assuming responsibility for the evaluation results report. <i>Note 1: For the implementation of the Mutual Acceptance Arrangement (MAA), Testing Laboratories may be part of the organization containing the OIML Issuing Authority or subcontracting Testing Laboratories of the OIML Issuing Authority.</i> <i>Note 2: Testing Laboratories may be a subcontractor of an inspection body.</i></p> <p>Partial Test Report Report issued by each Testing Laboratory, which includes results of tests and examinations that it performs.</p> |

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| | <p>OIML Partial Test Report Report, drawn up according to the Format specified in the relevant Recommendation for the applicable tests and examinations, issued by each Testing Laboratory, which includes results of tests and examinations it performs. <i>Note 1: In the event that the Issuing Authority has only one subcontracting Testing Laboratory, only one OIML Test Report may be issued.</i> <i>Note 2: In the event that the Issuing Authority performs the tests and examinations itself, the OIML Test Report may be directly combined with the OIML Evaluation Report (see below).</i></p> <p>Evaluation Report Report, issued by an Issuing Authority that includes the Partial Test Reports and assesses the conformity of the measuring instrument to stated requirements.</p> <p>OIML Evaluation Report Report, prepared according to the Test Report Format specified in the relevant Recommendation, that includes the OIML partial Test Reports and assesses the conformity of the type of measuring instrument to all the requirements in the relevant OIML Recommendation <i>Note 1: The OIML Evaluation Report is issued by the OIML Issuing Authority.</i> <i>Note 2: The OIML Evaluation Report is currently designated as “Test Report” in OIML B 3 and as “OIML Test Report” in OIML B 10-1.</i></p> <p>Additional Test Report Report issued by a Testing Laboratory that includes results of additional tests and examinations, behind those in the OIML Recommendation, accepted in the scope of a Declaration of Mutual Confidence (DoMC). <i>Note 1: Additional Test Reports are issued under the Mutual Acceptance Arrangement (MAA)</i> <i>Note 2: In the event that several Testing Laboratories are involved in the additional tests and examinations, each Testing Laboratory issues an Additional Test Report corresponding to those tests and examinations it performs.</i></p> <p>Complete Evaluation Report Report, issued by the OIML Issuing Authority, composed of the OIML Evaluation Report and of the additional Test Reports. <i>Note: Complete Evaluation Reports are issued under the Mutual Acceptance Arrangement (MAA)</i></p> <p>Abbreviations: MAA: Mutual Acceptance Arrangement DoMC: Declaration of Mutual Confidence CPR: Committee on Participation Review</p> |

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| <p>4 Management requirements</p> | |
| <p>4.1 Organization</p> | |
| <p>4.1.1 The laboratory or the organization of which it is part shall be an entity that can be held legally responsible.</p> | |
| <p>4.1.2 It is the responsibility of the laboratory to carry out its testing and calibration activities in such a way as to meet the requirements of this International Standard and to satisfy the needs of the customer, the regulatory authorities or organizations providing recognition.</p> | |
| <p>4.1.3 The management system shall cover work carried out in the laboratory's permanent facilities, at sites away from its permanent facilities, or in associated temporary or mobile facilities.</p> | <p>In particular, this requirement applies when Testing Laboratory personnel use a manufacturer's test facility to perform type approval tests and/or examinations.</p> |
| <p>4.1.4 If the laboratory is part of an organization performing activities other than testing and/or calibration, the responsibilities of key personnel in the organization that have an involvement or influence on the testing and/or calibration activities of the laboratory shall be defined in order to identify potential conflicts of interest.</p> <p><i>NOTE 1: Where a laboratory is part of a larger organisation, the organizational arrangements should be such that departments having conflicting interests, such as production, commercial marketing or financial do not adversely influence the laboratory's compliance with the requirements of this International Standard.</i></p> <p><i>NOTE 2: If the laboratory wishes to be recognized as a third-party laboratory, it should be able to demonstrate that it is impartial and that it and its personnel are free from any undue commercial, financial and other pressures which might influence their technical judgement. The third-party testing or calibration laboratory should not engage in any activities that may endanger the trust in its independence of judgement and integrity in relation to its testing or calibration activities.</i></p> | |
| <p>4.1.5 The laboratory shall:</p> <p>a) have managerial and technical personnel departures who, irrespective of other responsibilities, have the authority and resources needed to carry out their duties, including the implementation, maintenance and improvement of the management system, and to identify the occurrence of departures from the management system or from the procedures for performing tests and/or calibrations and to initiate actions to prevent or minimise such departures (see also 5.2);</p> | <p>d) Such policies shall be evaluated by Issuing Authorities when they designate Testing Laboratories.</p> |

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| <p>b) have arrangements to ensure that its management and personnel are free from any undue internal and external commercial, financial and other pressures that may adversely affect the quality of their work;</p> <p>c) have policies and procedures to ensure the protection of its customers' confidential information and proprietary rights, including procedures for protecting the electronic storage and transmission of results;</p> <p>d) have policies and procedures to avoid involvement in any activities that would diminish confidence in its competence, impartiality, judgement or operational integrity;</p> <p>e) define the organization and management structure of the laboratory, its place in any parent organization, and the relationships between quality management, technical operations, support services;</p> <p>f) specify the responsibility, authority and interrelationships of all personnel who manage, perform or verify work affecting the quality of the tests and/or calibrations;</p> <p>g) provide adequate supervision of testing and calibration staff, including trainees, by persons familiar with the methods and procedures, purpose of each test and/or calibration, and with the assessment of the test or calibration results;</p> <p>h) have technical management, which has overall responsibility for the technical operations and the provision of the resources needed to ensure the required quality of laboratory operations;</p> <p>i) appoint a member of staff as quality manager (however named) who, irrespective of other duties and responsibilities, shall have defined responsibility and authority for ensuring that the management system related to quality is implemented and followed at all times; the quality manager shall have direct access to the highest level of management at which decisions are made on laboratory policy or resources;</p> <p>j) appoint deputies for key managerial personnel (see Note);</p> <p>k) ensure that its personnel are aware of the relevance and importance of their activities and how they contribute to the achievement of the objectives of the management system.</p> <p><i>NOTE: Individuals may have more than one function and it may be impractical to appoint deputies for every function.</i></p> | |
| <p>4.1.6 Top management shall ensure that appropriate communication processes are established within the laboratory and that communication takes place regarding the effectiveness of the management system.</p> | |

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| <p>4.2 Management system</p> | |
| <p>4.2.1 The laboratory shall establish, implement and maintain a management system appropriate to the scope of its activities. The laboratory shall document its policies, systems, programmes, procedures and instructions to the extent necessary to assure the quality of the test and/or calibration results. The system's documentation shall be communicated to, understood by, available to, and implemented by the appropriate personnel.</p> | |
| <p>4.2.2 The laboratory's management system policies related to quality, including a quality policy statement, shall be defined in a quality manual (however named). The overall objectives shall be established, and shall be reviewed during management review. The quality policy statement shall be issued under the authority of top management. It shall include at least the following:</p> <p>a) the laboratory managements' commitment to good professional practice and to the quality of its testing and calibration in servicing its customers;</p> <p>b) the management's statement of the laboratory's standard of service;</p> <p>c) the purpose of the management system related to quality;</p> <p>d) a requirement that all personnel concerned with testing and calibration activities within the laboratory familiarize themselves with the quality documentation and implement the policies and procedures in their work; and</p> <p>e) the laboratory management's commitment to comply with this International Standard and to continually improve the effectiveness of the management system.</p> <p><i>NOTE:</i> <i>The quality policy statement should be concise and may include the requirement that tests and/or calibrations shall always be carried out in accordance with stated methods and customers' requirements. When the test and/or calibration laboratory is part of a larger organization, some quality policy elements may be in other documents.</i></p> | <p>b) In addition, the laboratory's policy shall include a statement that the testing process is in accordance with the instructions of the Issuing Authority. In the event of the implementation of the MAA, the statement shall specify that the testing process is in accordance with the relevant test procedures listed by the CPR.</p> |
| <p>4.2.3 Top management shall provide evidence of commitment to the development and implementation of the management system and to continually improving its effectiveness.</p> | |
| <p>4.2.4 Top management shall communicate to the organization the importance of meeting customer requirements as well as statutory and regulatory requirements.</p> | |
| <p>4.2.5 The quality manual shall include or make reference to the supporting procedures including technical procedures. It shall outline the structure of the documentation used in the management system.</p> | <p>In case of OIML Type Evaluations, technical procedures shall be in accordance with those defined in the relevant OIML Recommendation and, in case of the MAA implementation, with those defined in the DoMC for additional tests validated by the CPR.</p> |

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| <p>4.2.6 The roles and responsibilities of technical management and the quality manager, including their responsibility for ensuring compliance with this International Standard, shall be defined in the quality manual</p> | |
| <p>4.2.7 Top management shall ensure that the integrity of the management system is maintained when changes to the management system are planned and implemented.</p> | |
| <p>4.3 Document control</p> | |
| <p>4.3.1 General The laboratory shall establish and maintain procedures to control all documents that form part of its management system (internally generated and from external sources), such as regulations, standards, other normative documents, test and/or calibration methods, as well as drawings, software, specifications, instructions and manuals.</p> <p><i>NOTE 1: In this context “document” could be policy statements, procedures, specifications, calibration tables, charts, text books, posters, notices, memoranda, software, drawings, plans etc. These may be on various media whether hard copy or electronic, and they may be digital, analog, photographic or written.</i></p> <p><i>NOTE 2: The control of data related to testing and calibration is covered in clause 5.4.7. The control of records is covered in 4.13</i></p> | <p>The Testing Laboratory shall maintain updated documentation on the legal and contractual requirements applicable to its activity of evaluation testing and examinations (e.g. OIML Recommendations).</p> |
| <p>4.3.2 Document approval and issue</p> | |
| <p>4.3.2.1 All documents issued to personnel in the laboratory as part of the management system shall be reviewed and approved for use by authorized personnel prior to issue. A master list or an equivalent document control procedure identifying the current revision status and distribution of documents in the quality system shall be established and be readily available to preclude the use of invalid and/or obsolete documents.</p> | |
| <p>4.3.2.2 The procedure(s) adopted shall ensure that:</p> <ul style="list-style-type: none"> a) authorized editions of appropriate documents are available at all locations where operations essential to the effective functioning of the laboratory are performed; b) documents are periodically reviewed and, where necessary, revised to ensure continuing suitability and compliance with applicable requirements; c) invalid or obsolete documents are promptly removed from all points of issue or use, or otherwise assured against unintended use; d) obsolete documents retained for either legal or knowledge preservation purposes are suitably marked. | |

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| <p>4.3.2.3 Management system documents generated by the laboratory shall be uniquely identified. Such identification shall include the date of issue and/or revision identification, page numbering, the total number of pages or a mark to signify the end of the document, and the issuing authority(ies).</p> | |
| <p>4.3.3 Document changes</p> | |
| <p>4.3.3.1 Changes to documents shall be reviewed and approved by the same function that performed the original review unless specifically designated otherwise. The designated personnel shall have access to pertinent background information upon which to base their review and approval.</p> | |
| <p>4.3.3.2 Where practicable, the altered or new text shall be identified in the document or the appropriate attachments.</p> | |
| <p>4.3.3.3 If the laboratory's document control system allows for the amendment of documents by hand pending the re-issue of the documents, the procedures and authorities for such amendments shall be defined. Amendments shall be clearly marked, initialled and dated. A revised document shall be formally re-issued as soon as practicable.</p> | |
| <p>4.3.3.4 Procedures shall be established to describe how changes in documents maintained in computerized systems are made and controlled.</p> | |
| <p>4.4 Review of request, tender or contract</p> | |
| <p>4.4.1 The laboratory shall establish and maintain procedures for the review of requests, tenders or contracts. The policies and procedures for these reviews leading to a contract for testing and/or calibration shall ensure that:</p> <ul style="list-style-type: none"> a) the requirements, including the methods to be used, are adequately defined, documented and understood (see 5.4.2); b) the laboratory has the capability and resources to meet the requirements; c) the appropriate test and/or calibration method is selected and capable of meeting the customers' requirements (see 5.4.2). <p>Any differences between the request or tender and the contract shall be resolved before any work commences. Each contract shall be acceptable both to the laboratory and the customer. <i>NOTE 1: The request, tender and contract review should be conducted in a practical and efficient manner, and the effect of financial, legal and time schedule aspects should be taken into account. For internal customers, reviews of requests, tenders and contracts can be performed in a simplified way.</i></p> | <p>Each application shall lead to one contract only, covering all tests and examinations to be performed. This contract shall be signed by the Issuing Authority responsible for defining tests and examinations to be performed.</p> <p>On the basis of the contract signed by the Issuing Authority, each Testing Laboratory is responsible for reviewing the request considering tests and examinations it performs.</p> <ul style="list-style-type: none"> a) In the case of OIML Type Evaluation, examination and test procedures defined in the relevant OIML Recommendation and, in the event of the MAA implementation, in the DoMC for additional tests validated by the CPR, shall be used and understood. b) This requirement also applies if any test is performed by the Testing Laboratory which uses the manufacturer's test facilities. c) In the case of OIML Type Evaluation, requirements to be met are those defined in the relevant OIML Recommendation and, in the event of the MAA implementation, those defined in the additional requirements according to the scope of the DoMC. |

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| <p><i>NOTE 2: The review of capability should establish that the laboratory possesses the necessary physical, personnel and information resources, and that the laboratory's personnel have the skills and expertise necessary for the performance of the tests and/or calibrations in question. The review may also encompass results of earlier participation in interlaboratory comparisons or proficiency testing and/or the running of trial test or calibration programmes using samples or items of known value in order to determine uncertainties of measurement, limits of detection, confidence limits etc.</i></p> <p><i>NOTE 3: A contract may be any written or oral agreement to provide a customer with testing and/or calibration services.</i></p> | |
| <p>4.4.2 Records of reviews, including any significant changes, shall be maintained. Records shall also be maintained of pertinent discussions with a customer relating to the customer's requirements or the results of the work during the period of execution of the contract.</p> <p><i>NOTE: For review of routine and other simple tasks, the date and the identification (e.g. the initials) of the person in the laboratory responsible for carrying out the contracted work are considered adequate. For repetitive routine tasks, the review need be made only at the initial enquiry stage or on granting of the contract for on-going routine work performed under a general agreement with the customer, provided that the customer's requirements remain unchanged. For new, complex or advanced testing and/or calibration tasks a more comprehensive record should be maintained.</i></p> | |
| <p>4.4.3 The review shall also cover any work that is subcontracted by the laboratory.</p> | <p>Not applicable (see 4.5.1)</p> |
| <p>4.4.4 The customer shall be informed of any deviation from the contract.</p> | <p>Any deviation from the contract shall be submitted for approval to the Issuing Authority.</p> |
| <p>4.4.5 If a contract needs to be amended after work has commenced, the same contract review process shall be repeated and any amendments shall be communicated to all affected personnel.</p> | |
| <p>4.5 Subcontracting of tests and calibrations</p> | |
| <p>4.5.1 When a laboratory subcontracts work, whether because of unforeseen reasons (e.g. workload, need for further expertise or temporary incapacity) or on a continuing basis (e.g. through permanent subcontracting, agency or franchising arrangements), this work shall be placed with a competent subcontractor. A competent subcontractor is one that, for example, complies with this International Standard for the work in question.</p> | <p>Testing Laboratories involved in the type evaluation process are designated by the Issuing Authority.</p> <p>A Testing Laboratory is not authorized to subcontract any test and examination. Subcontracting in series is prohibited.</p> |

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| <p>4.5.2 The laboratory shall advise the customer of the arrangement in writing and, when appropriate, gain the approval of the customer preferably in writing.</p> | <p>Not applicable (see 4.5.1)</p> |
| <p>4.5.3 The laboratory is responsible to the customer for the subcontractor's work except in the case where the customer or a regulatory authority specifies which subcontractor is to be used.</p> | <p>Not applicable (see 4.5.1)</p> |
| <p>4.5.4 The laboratory shall maintain a register of all subcontractors that it uses for tests and/or calibrations and a record of the evidence of compliance with this International Standard for the work in question.</p> | <p>Not applicable (see 4.5.1). This responsibility belongs to the Issuing Authority when designating Testing Laboratories.</p> |
| <p>4.6 Purchasing services and supplies</p> | |
| <p>4.6.1 The laboratory shall have a policy and procedure(s) for the selection and purchasing of services and supplies it uses that affect the quality of the tests and/or calibrations. Procedures shall exist for the purchase, reception and storage of reagents and laboratory consumable materials relevant for the tests and calibrations.</p> | |
| <p>4.6.2 The laboratory shall ensure that purchased supplies and reagents and consumable materials that affect the quality of tests and/or calibrations are not used until they have been inspected or otherwise verified as complying with standard specifications or requirements defined in the methods for tests and/or calibrations concerned. These services and supplies used shall comply with specified requirements. Records of actions taken to check compliance shall be maintained.</p> | <p>In particular, services for calibration of test facilities are concerned to ensure appropriate uncertainties with the requirements.</p> |
| <p>4.6.3 Purchasing documents, for items affecting the quality of laboratory output shall contain data describing the services and supplies ordered. These purchasing documents shall be reviewed and approved for technical content prior to release.</p> <p><i>NOTE: The description may include type, class, grade, precise identification, specifications, drawings, inspection instructions, other technical data including approval of test results, the quality required and the management system standard under which they were made.</i></p> | |
| <p>4.6.4 The laboratory shall evaluate suppliers of critical consumables, supplies and services which affect the quality of testing and calibration, and shall maintain records of these evaluations and list those approved.</p> | |
| <p>4.7 Service to the customer</p> | |
| <p>4.7.1 The laboratory shall be willing to cooperate with customers or their representatives in clarifying the customer's request and in monitoring the laboratory's performance in relation to the work performed, provided that the laboratory ensures confidentiality to other customers.</p> | <p>Any clarification of the customer's request is under the responsibility of the Issuing Authority.</p> |

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| <p><i>NOTE 1: Such cooperation may include:</i></p> <p>a) providing the customer or the customer's representative reasonable access to relevant areas of the laboratory for the witnessing of tests and/or calibrations performed for the customer;</p> <p>b) preparation, packaging, and dispatch of test and/or calibration items needed by the customer for verification purposes.</p> <p><i>NOTE 2: Customers value maintenance of good communication, advice and guidance in technical matters, and opinions and interpretations based on results. Communication with the customer, especially in large assignments, should be maintained throughout the work. The laboratory should inform the customer of any delays or major deviations in the performance of the tests and/or calibrations.</i></p> | |
| <p>4.7.2 The laboratory shall seek feedback, both positive and negative, from its customers. The feedback shall be used and analysed to improve the management system, testing and calibration activities and customer service.</p> <p><i>NOTE: Examples of the types of feedback include customer satisfaction surveys and review of test or calibration reports with customers.</i></p> | |
| <p>4.8 Complaints</p> | |
| <p>The laboratory shall have a policy and procedure for the resolution of complaints received from customers or other parties. Records shall be maintained of all complaints and of the investigations and corrective actions taken by the laboratory (see also 4. 11).</p> | |
| <p>4.9 Control of nonconforming testing and/or calibration work</p> | |
| <p>4.9.1 The laboratory shall have a policy and procedures that shall be implemented when any aspect of its testing and/or calibration work, or the results of this work, do not conform to its own procedures or the agreed requirements of the customer. The policy and procedures shall ensure that:</p> <p>a) the responsibilities and authorities for the management of nonconforming work are designated and actions (including halting of work and withholding of test reports and calibration certificates, as necessary) are defined and taken when nonconforming work is identified;</p> <p>b) an evaluation of the significance of the nonconforming work is made;</p> <p>c) correction is taken immediately, together with any decision about the acceptability of the nonconforming work;</p> <p>d) where necessary, the customer is notified and work is recalled;</p> <p>e) the responsibility for authorizing the resumption of work is defined.</p> | <p>For decision to be taken, the Issuing Authority shall be informed of any nonconformance of test and examination results with the applicable requirements and of any major nonconformance in the implementation of test and examination procedures.</p> |

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| <p><i>NOTE: Identification of nonconforming work or problems with the management system or with testing and/or calibration activities can occur at various places within the management system and technical operations. Examples are customer complaints, quality control, instrument calibration, checking of consumable materials, staff observations or supervision, test report and calibration certificate checking, management reviews and internal or external audits.</i></p> | |
| <p>4.9.2 Where the evaluation indicates that the nonconforming work could recur or that there is doubt about the compliance of laboratory's operations with its own policies and procedures, the corrective action procedures given in 4.11 shall be promptly followed.</p> | |
| <p>4.10 Improvement</p> | |
| <p>The laboratory shall continually improve the effectiveness of its management system through the use of the quality policy, quality objectives, audit results, analysis of data, corrective and preventive actions and management review.</p> | |
| <p>4.11 Corrective action</p> | |
| <p>4.11.1 General The laboratory shall establish a policy and a procedure and shall designate appropriate authorities for implementing corrective action when nonconforming work or departures from the policies and procedures in the management system or technical operations have been identified.</p> <p><i>NOTE:</i> <i>A problem with the management system or with the technical operations of the laboratory may be identified through a variety of activities, such as control of nonconforming work, internal or external audits, management reviews, feedback from customers and staff observations.</i></p> | <p>This procedure shall include cooperation with the Issuing Authority in the event of nonconforming work.</p> |
| <p>4.11.2 Cause analysis The procedure for corrective action shall start with an investigation to determine the root cause(s) of the problem.</p> <p><i>NOTE: Cause analysis is the key and sometimes the most difficult part in the corrective action procedure. Often the root cause is not obvious and thus a careful analysis of all potential causes of the problem is required. Potential causes could include customer requirements, the samples, sample specifications, methods and procedures, staff skills and training, consumables, or equipment and its calibration.</i></p> | |

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| <p>4.11.3 Selection and implementation of corrective actions</p> <p>Where corrective action is needed, the laboratory shall identify the potential corrective actions. It shall select and implement the action(s) most likely to eliminate the problem and to prevent recurrence.</p> <p>Corrective actions shall be to a degree appropriate to the magnitude and the risk of the problem.</p> <p>The laboratory shall document and implement any required changes resulting from corrective action investigations.</p> | |
| <p>4.11.4 Monitoring of corrective actions</p> <p>The laboratory shall monitor the results to ensure that the actions taken have been effective.</p> | |
| <p>4.11.5 Additional audits</p> <p>Where the identification of nonconformities or departures casts doubts on the laboratory's compliance with its own policies and procedures, or on its compliance with this International Standard, the laboratory shall ensure that the appropriate areas of activity are audited in accordance with 4.14 as soon as possible.</p> <p><i>NOTE: Such additional audits often follow the implementation of the corrective actions to confirm their effectiveness. An additional audit should only be necessary when a serious issue or risk to the business is identified.</i></p> | <p>These additional audits may be internal or external. The Issuing Authority may require such an additional audit.</p> |
| <p>4.12 Preventive action</p> | |
| <p>4.12.1 Needed improvements and potential sources of nonconformities, either technical or concerning the management system, shall be identified. When improvement opportunities are identified or if preventive action is required, action plans shall be developed, implemented and monitored to reduce the likelihood of the occurrence of such nonconformities and to take advantage of the opportunities for improvement.</p> | |
| <p>4.12.2 Procedures for preventive actions shall include the initiation of such actions and the application of controls to ensure that they are effective.</p> <p><i>NOTE 1: Preventive action is a pro-active process to identify opportunities for improvement rather than a reaction to the identification of problems or complaints.</i></p> <p><i>NOTE 2: Apart from the review of the operational procedures, the preventive action might involve analysis of data, including trend and risk analyses and proficiency-testing results.</i></p> | |

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| <p>4.13 Control of records</p> | |
| <p>4.-13.1 General</p> | |
| <p>4.13.1.1 The laboratory shall establish and maintain procedures for identification, collection, indexing, access, filing, storage, maintenance and disposal of quality and technical records. Quality records shall include reports from internal audits and management reviews as well as corrective and preventive actions.</p> | |
| <p>4.13.1.2 All records shall be legible and shall be stored and retained in such a way that they are readily retrievable in facilities that provide a suitable environment to prevent damage or deterioration and to prevent loss. Retention times of records shall be established.</p> <p><i>NOTE: Records may be in any media, such as hard copy or electronic media.</i></p> | <p>In the case of OIML Type Evaluation, technical records related to type evaluation tests and examinations shall be maintained available as long as the OIML Certificate remains registered.</p> |
| <p>4.13.1.3 All records shall be held secure and in confidence.</p> | |
| <p>4.13.1.4 The laboratory shall have procedures to protect and back up records stored electronically and to prevent unauthorized access to or amendment of these records.</p> | |
| <p>4.13.2 Technical records</p> | |
| <p>4.13.2.1 The laboratory shall retain original observations, derived data and sufficient information to establish an audit trail, calibration records, staff records and a copy of each test report or calibration certificate issued, for a defined period. The records for each test or calibration shall contain sufficient information to facilitate, if possible, identification of factors affecting the uncertainty and to enable the test or calibration to be repeated under conditions as close as possible to the original. The records shall include the identity of personnel responsible for the sampling, performance of each test and/or calibration and checking of results.</p> <p><i>NOTE 1: In certain fields it may be impossible or impractical to retain records of all original observations.</i></p> <p><i>NOTE 2: Technical records are accumulations of data (see 5.4.7) and information which result from carrying out tests and/or calibrations and which indicate whether specified quality or process parameters are achieved. They may include forms, contracts, work sheets, work books, check sheets, work notes, control graphs, external and internal test reports and calibration certificates, customers' notes, papers and feedback.</i></p> | |
| <p>4.13.2.2 Observations, data and calculations shall be recorded at the time they are made and shall be identifiable to the specific task.</p> | |

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| <p>4.13.2.3 When mistakes occur in records, each mistake shall be crossed out, not erased, made illegible or deleted, and the correct value entered alongside. All such alterations to records shall be signed or initialled by the person making the correction. In the case of records stored electronically, equivalent measures shall be taken to avoid loss or change of original data.</p> | |
| <p>4.14 Internal audits</p> | |
| <p>4.14.1 The laboratory shall periodically, and in accordance with a predetermined schedule and procedure, conduct internal audits of its activities to verify that its operations continue to comply with the requirements of the management system and this International Standard. The internal audit programme shall address all elements of the management system, including the testing and/or calibration activities. It is the responsibility of the quality manager to plan and organize audits as required by the schedule and requested by management. Such audits shall be carried out by trained and qualified personnel who are, wherever resources permit, independent of the activity to be audited.</p> <p><i>NOTE: The cycle for internal auditing should normally be completed in one year.</i></p> | |
| <p>4.14.2 When audit findings cast doubt on the effectiveness of the operations or on the correctness or validity of the laboratory's test or calibration results, the laboratory shall take timely corrective action, and shall notify customers in writing, if investigations show that the laboratory results may have been affected.</p> | |
| <p>4.14.3 The area of activity audited, the audit findings and corrective actions that arise from them shall be recorded.</p> | |
| <p>4.14.4 Follow up audit activities shall verify and record the implementation and effectiveness of the corrective action taken.</p> | |
| <p>4.15 Management reviews</p> | |
| <p>4.15.1 In accordance with a predetermined schedule and procedure, the laboratory's top management shall periodically conduct a review of the laboratory's management system and testing and/or calibration activities to ensure their continuing suitability and effectiveness, and to introduce necessary changes or improvements. The review shall take account of:</p> <ul style="list-style-type: none"> - the suitability of policies and procedures; - reports from managerial and supervisory personnel; - the outcome of recent internal audits; - corrective and preventive actions; - assessments by external bodies; - the results of interlaboratory comparisons or proficiency tests; | <p>Fifth bullet: Audits performed on Testing Laboratories by the external Issuing Authority, carried out under the MAA are considered as assessments by external bodies and their results should be taken into account in the management review of the Testing Laboratory.</p> |

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| <ul style="list-style-type: none"> - changes in the volume and type of the work; - customer feedback; - complaints; - recommendations for improvement; - other relevant factors, such as quality control activities, resources and staff training. <p><i>NOTE 1: A typical period for conducting a management review is once every 12 months.</i></p> <p><i>NOTE 2: Results should feed into the laboratory planning system and should include the goals, objectives and action plans for the coming year.</i></p> <p><i>NOTE 3: A management review includes consideration of related subjects at regular management meetings.</i></p> | |
| <p>4.15.2 Findings from management reviews and the actions that arise from them shall be recorded. The management shall ensure that those actions are carried out within an appropriate and agreed timescale.</p> | <p>The findings from management reviews shall be submitted to the Issuing Authority.</p> |
| <p>5 Technical requirements</p> | |
| <p>5.1 General</p> | |
| <p>5.1.1 Many factors determine the correctness and reliability of the tests and/or calibrations performed by a laboratory. These factors include contributions from:</p> <ul style="list-style-type: none"> - human factors (5.2); - accommodation and environmental conditions (5.3); - test and calibration methods and method validation (5.4); - equipment (5.5); - measurement traceability (5.6); - sampling (5.7); - the handling of test and calibration items (5.8). | |
| <p>5.1.2 The extent to which the factors contribute to the total uncertainty of measurement differs considerably between (types of) tests and between (types of) calibrations. The laboratory shall take account of these factors in developing test and calibration methods and procedures, in the training and qualification of personnel, and in the selection and calibration of the equipment it uses.</p> | |
| <p>5.2 Personnel</p> | |
| <p>5.2.1 The laboratory management shall ensure the competence of all who operate specific equipment, perform tests and/or calibrations, evaluate results, and sign test reports and calibration certificates. When using staff who are undergoing training, appropriate supervision shall be provided. Personnel performing specific tasks shall be qualified on the basis of appropriate education, training, experience and/or demonstrated skills, as required.</p> | <p>Training methods for the personnel should include participation in international work in the field of legal metrology (e.g. development of OIML Publications). This includes work for OIML performed at national level.</p> |

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| <p><i>NOTE 1: In some technical areas (e.g. non-destructive testing) it may be required that the personnel performing certain tasks hold personnel certification. The laboratory is responsible for fulfilling specified personnel certification requirements. The requirements for personnel certification might be regulatory, included in the standards for the specific technical field, or required by the customer. NOTE 2: The personnel responsible for the opinions and interpretation included in test reports should, in addition to the appropriate qualifications, training, experience and satisfactory knowledge of the testing carried out, also have:</i></p> <ul style="list-style-type: none"> - <i>relevant knowledge of the technology used for the manufacturing of the items, materials, products, etc. tested, or the way they are used or intended to be used, and of the defects or degradations which may occur during or in service;</i> - <i>knowledge of the general requirements expressed in the legislation and standards; and</i> - <i>an understanding of the significance of deviations found with regard to the normal use of the items, materials, products, etc. concerned.</i> | <p><i>NOTE 2:</i> In particular, this note is applicable to the personnel responsible for:</p> <ul style="list-style-type: none"> - the examinations (which are part of the evaluation); - the statement of the compliance of tests results with requirements (see 5.10.5). |
| <p>5.2.2 The management of the laboratory shall formulate the goals with respect to the education, training and skills of the laboratory personnel. The laboratory shall have a policy and procedures for identifying training needs and providing training of personnel. The training programme shall be relevant to the present and anticipated tasks of the laboratory. The effectiveness of the training actions taken shall be evaluated.</p> | <p>The personnel in charge of type evaluation testing and examinations shall be aware of the following:</p> <ul style="list-style-type: none"> - relevant OIML Publications; - these guidelines; - applicable Declarations of Mutual Confidence; <p>OIML Publication D 14 gives guidelines for training of legal metrology personnel.</p> |
| <p>5.2.3 The laboratory shall use personnel who are employed by, or under contract to, the laboratory. Where contracted and additional technical and key support personnel are used, the laboratory shall ensure that such personnel are supervised and competent and that they work in accordance with the laboratory's management system.</p> | |
| <p>5.2.4 The laboratory shall maintain current job descriptions for managerial, technical and key support personnel involved in tests and/or calibrations.</p> <p><i>NOTE: Job descriptions can be defined in many ways. As a minimum, the following should be defined:</i></p> <ul style="list-style-type: none"> - <i>the responsibilities with respect to performing tests and/or calibrations;</i> - <i>the responsibilities with respect to the planning of tests and/or calibrations and evaluation of results;</i> - <i>the responsibilities for reporting opinions and interpretations;</i> - <i>the responsibilities with respect to method modification and development and validation of new methods;</i> | |

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| <ul style="list-style-type: none"> - <i>expertise and experience required;</i> - <i>qualifications and training programmes;</i> - <i>managerial duties.</i> | |
| <p>5.2.5 The management shall authorize specific personnel to perform particular types of sampling, test and/or calibration, to issue test reports and calibration certificates, to give opinions and interpretations and to operate particular types of equipment. The laboratory shall maintain records of the relevant authorization(s), competence, educational and professional qualifications, training, skills and experience of all technical personnel, including contracted personnel. This information shall be readily available and shall include the date on which authorization and/or competence is confirmed.</p> | <p>Tests and examinations shall be performed by personnel having appropriate competence. This competence shall be evaluated and validated by the Testing Laboratory's technical management (see 4.1.5 f)).</p> <p>A list shall be kept up to date, indicating for each category of measuring instruments:</p> <ul style="list-style-type: none"> - personnel qualified to carry out tests and/or examinations; - personnel qualified to give an opinion on the statement of compliance/non compliance of the results with requirements; - personnel responsible for training; - managerial personnel responsible for validating technical work. <p>Personnel in the process of being qualified shall be in charge only of simple or well described activities. They can participate in, but not be responsible for, testing.</p> |
| <p>5.3 Accommodation and environmental conditions</p> | |
| <p>5.3.1 Laboratory facilities for testing and/or calibration, including but not limited to energy sources, lighting and environmental conditions, shall be such as to facilitate correct performance of the tests and/or calibrations.</p> <p>The laboratory shall ensure that the environmental conditions do not invalidate the results or adversely affect the required quality of any measurement. Particular care shall be taken when sampling tests and/or calibrations are undertaken at sites other than a permanent laboratory facility. The technical requirements for accommodation and environmental conditions that can affect the results of tests and calibrations shall be documented.</p> | <p>Second paragraph: Standards and test equipment shall be used in the same environmental conditions as those of their calibration. Otherwise, appropriate corrections or new uncertainty calculations shall be performed. In the event that this is not possible, test results shall not be valid. The policy related to this matter shall be documented.</p> |
| <p>5.3.2 The laboratory shall monitor, control and record environmental conditions as required by the relevant specifications, methods and procedures or where they influence the quality of the results. Due attention shall be paid, for example, to biological sterility, dust, electromagnetic disturbances, radiation, humidity, electrical supply, temperature, and sound and vibration levels, as appropriate to the technical activities concerned. Tests and calibrations shall be stopped when the environmental conditions jeopardize the results of the tests and/or calibrations.</p> | <p>In particular, information related to rain, wind etc. shall be recorded in the case of tests performed outdoors.</p> |
| <p>5.3.3 There shall be effective separation between neighbouring areas in which there are incompatible activities. Measures shall be taken to prevent cross-contamination.</p> | |

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| <p>5.3.4 Access to and use of areas affecting the quality of the tests and/or calibrations shall be controlled. The laboratory shall determine the extent of control based on its particular circumstances.</p> | |
| <p>5.3.5 Measures shall be taken to ensure good housekeeping in the laboratory. Special procedures shall be prepared where necessary.</p> | |
| <p>5.4 Test and calibration methods and method validation</p> | |
| <p>5.4.1 General The laboratory shall use appropriate methods and procedures for all tests and/or calibrations within its scope. These include sampling, handling, transport, storage and preparation of items to be tested and/or calibrated, and, where appropriate, an estimation of the measurement uncertainty as well as statistical techniques for analysis of test and/or calibration data.</p> <p>The laboratory shall have instructions on the use and operation of all relevant equipment, and on the handling and preparation of items for testing and/or calibration, or both, where the absence of such instructions could jeopardize the results of tests and/or calibrations. All instructions, standards, manuals and reference data relevant to the work of the laboratory shall be kept up to date and shall be made readily available to personnel (see 4.3). Deviation from test and calibration methods shall occur only if the deviation has been documented, technically justified, authorized and accepted by the customer.</p> <p><i>NOTE : International, regional or national standards or other recognized specifications that contain sufficient and concise information to perform the tests and/or calibrations, do not need to be supplemented or rewritten as internal procedures if these standards are written in a way that they can be used as published by the operating staff in a laboratory. It may be necessary to provide additional documentation for optional steps in the method or additional details.</i></p> | <p>These procedures shall ensure that the Testing Laboratory verifies that:</p> <ul style="list-style-type: none"> - the sample(s) to be examined and/or tested are those validated by the Issuing Authority, - the configuration of the sample(s) to be examined and/or tested is the original one. <p>Selection of instruments to be tested amongst a family is considered as sampling.</p> <p>These procedures shall indicate how adjustments and modifications of the sample(s) authorized by the Issuing Authority are taken into account during tests and examinations.</p> <p>These requirements also apply when tests are performed with the manufacturer's test facilities.</p> |

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| <p>5.4.2 Selection of methods</p> <p>The laboratory shall use test and/or calibration methods, including methods for sampling, which meet the needs of the customer and which are appropriate for the tests and/or calibrations it undertakes. Methods published in international, regional or national standards shall preferably be used. The laboratory shall ensure that it uses the latest valid edition of a standard unless it is not appropriate or possible to do so. When necessary, the standard shall be supplemented with additional details to ensure consistent application. When the customer does not specify the method to be used, the laboratory shall select appropriate methods that have been published either in international, regional or national standards, or by reputable technical organizations, or in relevant scientific texts or journals, or as specified by the manufacturer of the equipment. Laboratory-developed methods or methods adopted by the laboratory may also be used if they are appropriate for the intended use and if they are validated. The customer shall be informed as to the method chosen. The laboratory shall confirm that it can properly operate standard methods before introducing the tests or calibrations. If the standard method changes, the confirmation shall be repeated.</p> <p>The laboratory shall inform the customer when the method proposed by the customer is considered to be inappropriate or out-of-date.</p> | <p>See 4.2.2</p> <p>In the case of OIML Type Evaluation, the test and examination procedures shall conform to those defined in the applicable OIML Recommendation and, if applicable, to the additional procedures validated by the DoMC in case of the MAA implementation.</p> |
| <p>5.4.3 Laboratory-developed methods</p> <p>The introduction of test and calibration methods developed by the laboratory for its own use shall be a planned activity and be assigned to qualified personnel equipped with adequate resources.</p> <p>Plans shall be updated as development proceeds and effective communication amongst all personnel involved shall be ensured.</p> | <p>Not applicable (see 5.4.2).</p> |
| <p>5.4.4 Non standardized methods</p> <p>When it is necessary to use methods not covered by standard methods, these shall be subject to agreement with the customer and shall include a clear specification of the customers' requirements and the purpose of the test and/or calibration. The method developed shall have been validated appropriately before use.</p> <p><i>NOTE: For new test and/or calibration methods, procedures should be developed prior to the tests and/or calibrations being performed and should contain at least the following information:</i></p> <ul style="list-style-type: none"> <i>a) appropriate indication;</i> <i>b) scope;</i> <i>c) description of the type of item to be tested or calibrated;</i> <i>d) parameters or quantities to be determined;</i> | <p>Not applicable (see 5.4.2).</p> <p>Test methods validated by the CPR are considered as standardized methods.</p> |

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| <p>e) apparatus and equipment, including technical requirement;</p> <p>f) reference standards and reference materials required;</p> <p>g) environmental conditions required and any stabilization period needed;</p> <p>h) description of the procedure, including</p> <ul style="list-style-type: none"> - affixing of identification marks, handling, transporting, storing and preparing of items, - checks to be made before the work is started, - checks that the equipment is working properly and, where required, calibration and adjustment of the equipment before each use, - the method of recording the observations and results, - any safety measures to be observed; <p>i) criteria and/or requirements for approval/rejection;</p> <p>j) data to be recorded and method of analysis and presentation;</p> <p>k) the uncertainty or the procedure for estimating uncertainty.</p> | |
| <p>5.4.5 Validation of methods</p> | |
| <p>5.4.5.1 Validation is the confirmation by examination and the provision of objective evidence that the particular requirements for a specific intended use are fulfilled.</p> | <p>Validation of methods is under the responsibility of higher authorities (CIML for OIML Recommendations and CPR for additional tests).</p> |
| <p>5.4.5.2 The laboratory shall validate non-standard methods, laboratory designed/developed methods, standard methods used outside their intended scope and amplifications and modifications of standard methods to confirm that the methods are fit for the intended use. The validation shall be as extensive as is necessary to meet the needs in the given application or field of application. The laboratory shall record the results obtained, the procedure used for the validation, and a statement as to whether the method is fit for the intended use.</p> <p><i>NOTE 1: Validation may include procedure for sampling, handling and transportation.</i></p> <p><i>NOTE 2: The techniques used for the determination of the performance of a method should be one of, or a combination of, the following:</i></p> <ul style="list-style-type: none"> - calibration using reference standards or reference materials; - comparison of results achieved with other methods; - interlaboratory comparisons; - systematic assessment of the factors influencing the result; - assessment of the uncertainty of the results based on scientific understanding of the theoretical principles of the method and practical experience. | <p>Not applicable (see 5.4.2 and 5.4.5.1).</p> |

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| <p><i>NOTE 3: When some changes are made in the validated non-standard methods, the influence of such changes should be documented and, if appropriate, a new validation should be carried out.</i></p> | |
| <p>5.4.5.3 The range and accuracy of the values obtainable from validated methods (e.g. the uncertainty of the results, detection limit, selectivity of the method, linearity, limit of repeatability and/or reproducibility, robustness against external influences and/or cross-sensitivity against interference from the matrix of the sample/test object), as assessed for the intended use, shall be relevant to the customers' needs.</p> <p><i>NOTE 1: Validation includes the specification of the requirements, determination of the characteristics of the methods, a check that the requirements can be fulfilled by using the method, and a statement on the validity.</i></p> <p><i>NOTE 2: As method-development proceeds, regular review should be carried out to verify that the needs of the customer are still being fulfilled. Any change in requirements requiring modifications to the development plan shall be approved and authorized.</i><i>NOTE 3: Validation is always a balance between costs, risks and technical possibilities. There are many cases in which the range and uncertainty of the values (e.g. accuracy, detection limit, selectivity, linearity, repeatability, reproducibility, robustness and cross-sensitivity) can only be given in a simplified way due to lack of information.</i></p> | <p>See 5.4.5.1.</p> |
| <p>5.4.6 Estimation of uncertainty of measurement</p> | |
| <p>5.4.6.1 A calibration laboratory, or a testing laboratory performing its own calibrations, shall have and shall apply a procedure to estimate the uncertainty of measurement for all calibrations and types of calibrations.</p> | |
| <p>5.4.6.2 Testing laboratories shall have and shall apply procedures for estimating uncertainty of measurement. In certain cases the nature of the test method may preclude rigorous, metrologically and statistically valid, calculation of uncertainty of measurement. In these cases the laboratory shall at least attempt to identify all the components of uncertainty and make a reasonable estimation, and shall ensure that the form of reporting of the result does not give a wrong impression of the uncertainty. Reasonable estimation shall be based on knowledge of the performance of the method and on the measurement scope and shall make use of, for example, previous experience and validation data.</p> | <p>Testing laboratories shall include in these procedures the confirmation of the acceptable uncertainties of measurements with the requirements of the relevant OIML Recommendation.</p> <p><i>NOTE: If the laboratory is accredited by a recognized accreditation body, the adequacy of the uncertainty calculations is deemed to be demonstrated if type evaluation testing falls under the scope of the accreditation. In that case, according to OIML B 10, no peer assessment is required.</i></p> |

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| <p><i>NOTE 1: The degree of rigor needed in an estimation of uncertainty of measurement depends on factors such as:</i></p> <ul style="list-style-type: none"> - <i>the requirements of the test method;</i> - <i>the requirements of the customer;</i> - <i>the existence of narrow limits on which decisions on conformity to a specification are based.</i> <p><i>NOTE 2: In those cases where a well- recognised test method specifies limits to the values of the major sources of uncertainty of measurement and specifies the form of presentation of calculated results, the laboratory is considered to have satisfied this clause by following the test method and reporting instructions (see 5.10).</i></p> | |
| <p>5.4.6.3 When estimating the uncertainty of measurement, all uncertainty components which are of importance in the given situation shall be taken into account using appropriate methods of analysis.</p> <p><i>NOTE 1: Sources contributing to the uncertainty include, but are not necessarily limited to, the reference standards and reference materials used, methods and equipment used, environmental conditions, properties and condition of the item being tested or calibrated, and the operator.</i></p> <p><i>NOTE 2: The predicted long-term behaviour of the tested and/or calibrated item is normally not taken into account when estimating the measurement uncertainty.</i></p> <p><i>NOTE 3: For further information, see ISO 5725 and the Guide to the Expression of Uncertainty in Measurement (see Bibliography).</i></p> | |
| <p>5.4.7 Control of data</p> | |
| <p>5.4.7.1 Calculations and data transfers shall be subject to appropriate checks in a systematic manner.</p> | |
| <p>5.4.7.2 When computers or automated equipment are used for the acquisition, processing, recording, reporting, storage or retrieval of test or calibration data, the laboratory shall ensure that:</p> <ol style="list-style-type: none"> a) computer software developed by the user is documented in sufficient detail and suitably validated as being adequate for use; b) procedures are established and implemented for protecting the data ; such procedures shall include, but not be limited to, integrity and confidentiality of data entry or collection, data storage, data transmission and data processing; c) computers and automated equipment are maintained to ensure proper functioning and are provided with the environmental and operating conditions necessary to maintain the integrity of test and calibration data. | |

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| <p><i>NOTE: Commercial off-the-shelf software (e.g. wordprocessing, database and statistical programmes) in general use within their designed application range may be considered to be sufficiently validated. However, laboratory software configuration/modifications should be validated as in 5.4.7.2a).</i></p> | |
| <p>5.5 Equipment</p> | |
| <p>5.5.1 The laboratory shall be furnished with all items of sampling, measurement and test equipment required for the correct performance of the tests and/or calibrations (including sampling, preparation of test and/or calibration items, processing and analysis of test and/or calibration data). In those cases where the laboratory needs to use equipment outside its permanent control, it shall ensure that the requirements of this International Standard are met.</p> | <p>The last sentence of the requirement applies when the Testing Laboratory uses the manufacturer's test facilities.</p> |
| <p>5.5.2 Equipment and its software used for testing, calibration and sampling shall be capable of achieving the accuracy required and shall comply with specifications relevant to the tests and/or calibrations concerned. Calibration programmes shall be established for key quantities or values of the instruments where these properties have a significant effect on the results. Before being placed into service, equipment (including that used for sampling) shall be calibrated or checked to establish that it meets the laboratory's specification requirements and complies with the relevant standard specifications. It shall be checked and/or calibrated before use (see 5.6).</p> | <p>See 5.4.6.2.</p> |
| <p>5.5.3 Equipment shall be operated by authorized personnel. Up-to-date instructions on the use and maintenance of equipment (including any relevant manuals provided by the manufacturer of the equipment) shall be readily available for use by the appropriate laboratory personnel.</p> | |
| <p>5.5.4 Each item of equipment and its software used for testing and calibration and significant to the result shall, when practicable, be uniquely identified.</p> | |
| <p>5.5.5 Records shall be maintained of each item of equipment and its software significant to the tests and/or calibrations performed. The records shall include at least the following:</p> <ul style="list-style-type: none"> a) identity of the item of equipment and its software; b) the manufacturer's name, type identification, and serial number or other unique identification; c) checks that equipment complies with the specification (see 5.5.2); d) the current location, where appropriate; e) the manufacturer's instructions, if available, or reference to their location; f) dates, results and copies of reports and certificates of all calibrations, adjustments, acceptance criteria, and the due date of next calibration; | |

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| <p>g) the maintenance plan, where appropriate, and maintenance carried out to date ; h) any damage, malfunction, modification or repair to the equipment.</p> | |
| <p>5.5.6 The laboratory shall have procedures for safe handling, transport, storage, use and planned maintenance of measuring equipment to ensure proper functioning and in order to prevent contamination or deterioration.</p> <p><i>NOTE: Additional procedures may be necessary when measuring equipment is used outside the permanent laboratory for tests, calibrations or sampling.</i></p> | |
| <p>5.5.7 Equipment that has been subjected to overloading or mishandling, gives suspect results, or has been shown to be defective or outside specified limits, shall be taken out of service. It shall be isolated to prevent its use or clearly labelled or marked as being out of service until it has been repaired and shown by calibration or test to perform correctly. The laboratory shall examine the effect of the defect or departure from specified limits on previous tests and/or calibrations and shall institute the "Control of nonconforming work" procedure (see 4.9).</p> | |
| <p>5.5.8 Whenever practicable, all equipment under the control of the laboratory, and requiring calibration shall be labelled, coded or otherwise identified to indicate the status of calibration including, the date when last calibrated and the date or expiration criteria when recalibration is due.</p> | |
| <p>5.5.9 When, for whatever reason, equipment goes outside the direct control of the laboratory, the laboratory shall ensure that the function and calibration status of the equipment are checked and shown to be satisfactory before the equipment is returned to service.</p> | <p>Also applies when the Testing Laboratory uses the manufacturer's test facilities.</p> |
| <p>5.5.10 When intermediate checks are needed to maintain confidence in the calibration status of the equipment, these checks shall be carried out according to a defined procedure.</p> | |
| <p>5.5.11 Where calibrations give rise to a set of correction factors, the laboratory shall have procedures to ensure that copies (e.g. in computer software) are correctly updated.</p> | |
| <p>5.5.12 Test and calibration equipment, including both hardware and software, shall be safeguarded from adjustments which would invalidate the test and/or calibration results.</p> | |

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| <p>5.6 Measurement traceability</p> | |
| <p>5.6.1 General All equipment used for tests and/or calibrations, including equipment for subsidiary measurements (e.g. for environmental conditions) having a significant effect on the accuracy or validity of the result of the test, calibration or sampling shall be calibrated before being put into service. The laboratory shall have an established programme and procedure for the calibration of its equipment.</p> <p><i>NOTE: Such a programme should include a system for selecting, using, calibrating, checking, controlling and maintaining measurement standards, reference materials used as measurement standards, and measuring and test equipment used to perform tests and calibrations.</i></p> | |
| <p>5.6.2 Specific requirements</p> | |
| <p>5.6.2.1 Calibration</p> | |
| <p>5.6.2.1.1 For calibration laboratories, the programme for calibration of equipment shall be designed and operated so as to ensure that calibrations and measurements made by the laboratory are traceable to the International System of (SI) (Système international d'unités).</p> <p>A calibration laboratory establishes traceability of its own measurement standards and measuring instruments to SI by means of an unbroken chain of calibrations or comparisons linking them to relevant primary standards of the SI units of measurement. The link to SI units may be achieved by reference to national measurement standards. National measurement standards may be primary standards, which are primary realizations of the SI units or agreed representations of SI units based on fundamental physical constants, or they may be secondary standards which are standards calibrated by another national metrology institute. When using external calibration services, traceability of measurement shall be assured by the use of calibration services from laboratories that can demonstrate competence, measurement capability and traceability. The calibration certificates issued by these laboratories shall contain the measurement results, including the measurement uncertainty and/or a statement of compliance with an identified metrological specification (see also 5.10.4.2).</p> <p><i>NOTE 1: Calibration laboratories fulfilling the requirements of this International Standard are considered to be competent. A calibration certificate bearing an accreditation body logo from a calibration laboratory accredited to this International Standard, for the calibration concerned, is sufficient evidence of traceability of the calibration data reported.</i></p> | <p>See 5.6.2.2.1 5.6.2.1 of ISO/IEC 17025 is related to Calibration Laboratories which are not the object of this Guide.</p> |

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| <p><i>NOTE 2: Traceability to SI units of measurement may be achieved by reference to an appropriate primary standard (see VIM: 1993, 6.4) or by reference to a natural constant, the value of which in terms of the relevant SI unit is known and recommended by the General Conference of Weights and Measures (CGPM) and the International Committee for Weights and Measures (CIPM).</i></p> <p><i>NOTE 3: Calibration laboratories that maintain their own primary standard or representation of SI units based on fundamental physical constants can claim traceability to the SI system only after these standards have been compared, directly or indirectly, with other similar standards of a national metrology institute.</i></p> <p><i>NOTE 4: The term “identified metrological specification” means that it must be clear from the calibration certificate which specification the measurements have been compared with, by including the specification or by giving an unambiguous reference to the specification.</i></p> <p><i>NOTE 5: When the terms “international standard” or “national standard” are used in connection with traceability, it is assumed that these standards fulfil the properties of primary standards for the realization of SI units.</i></p> <p><i>NOTE 6: Traceability to national measurement standards does not necessarily require the use of the national metrology institute of the country in which the laboratory is located.</i></p> <p><i>NOTE 7: If a calibration laboratory wishes or needs to obtain traceability from a national metrology institute other than in its own country, this laboratory should select a national metrology institute that actively participates in the activities of BIPM either directly or through regional groups.</i></p> <p><i>NOTE 8: The unbroken chain of calibrations or comparisons may be achieved in several steps carried out by different laboratories that can demonstrate traceability.</i></p> | |
| <p>5.6.2.1.2 There are certain calibrations that currently cannot be strictly made in SI units. In these cases calibration shall provide confidence in measurements by establishing traceability to appropriate measurement standards such as:</p> <ul style="list-style-type: none"> - the use of certified reference materials provided by a competent supplier to give a reliable physical or chemical characterization of a material; - the use of specified methods and/or consensus standards that are clearly described and agreed by all parties concerned. | |

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| <p>Participation in a suitable programme of interlaboratory comparisons is required where possible.</p> | |
| <p>5.6.2.2 Testing</p> | |
| <p>5.6.2.2.1 For testing laboratories, the requirements given in 5.6.2.1 apply for measuring and test equipment with measuring functions used, unless it has been established that the associated contribution from the calibration contributes little to the total uncertainty of the test result. When this situation arises, the laboratory shall ensure that the equipment used can provide the uncertainty of measurement needed.</p> <p><i>NOTE: The extent to which the requirements in 5.6.2.1 should be followed depends on the relative contribution of the calibration uncertainty to the total uncertainty. If calibration is the dominant factor, the requirements should be strictly followed.</i></p> | <p>Any external calibration shall be performed by an accredited calibration laboratory or by a national metrology institute.</p> <p>For the implementation of the MAA, any external calibration shall be performed by a calibration laboratory accredited by an accreditation body that is a full Member of ILAC.</p> <p>In the event that the acceptance criteria is either stability or based on a fault determination, complete traceability may not be necessary. In such a case the repeatability of the test equipment is much more important than the true value as indicated by the equipment, since the requirement is based on differences of errors. To this end, a calibration certificate from an accredited laboratory may not be necessary.</p> |
| <p>5.6.2.2.2 Where traceability of measurements to SI units is not possible and/or not relevant, the same requirements for traceability to, for example, certified reference materials, agreed methods and/or consensus standards, are required as for calibration laboratories (see 5.6.2.1.2).</p> | |
| <p>5.6.3 Reference standards and reference materials</p> | |
| <p>5.6.3.1 Reference standards</p> <p>The laboratory shall have a programme and procedure for the calibration of its reference standards. Reference standards shall be calibrated by a body that can provide traceability as described in 5.6.2.1. Such reference standards of measurement held by the laboratory shall be used for calibration only and for no other purpose, unless it can be shown that their performance as reference standards would not be invalidated. Reference standards shall be calibrated before and after any adjustment.</p> | |
| <p>5.6.3.2 Reference materials</p> <p>Reference materials shall, where possible, be traceable to SI units of measurement, or to certified reference materials. Internal reference materials shall be checked as far as is technically and economically practicable.</p> | |
| <p>5.6.3.3 Intermediate checks</p> <p>Checks needed to maintain confidence in the calibration status of reference, primary, transfer or working standards and reference materials shall be carried out according to defined procedures and schedules.</p> | |

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| <p>5.6.3.4 Transport and storage</p> <p>The laboratory shall have procedures for safe handling, transport, storage and use of reference standards and reference materials in order to prevent contamination or deterioration and in order to protect their integrity.</p> <p><i>NOTE: Additional procedures may be necessary when reference standards and reference materials are used outside the permanent laboratory for tests, calibrations or sampling.</i></p> | |
| <p>5.7 Sampling</p> | |
| <p>5.7.1 The laboratory shall have a sampling plan and procedures for sampling when it carries out sampling of substances, materials or products for subsequent testing or calibration. The sampling plan as well as the sampling procedure shall be available at the location where sampling is undertaken. Sampling plans shall, whenever reasonable, be based on appropriate statistical methods. The sampling process shall address the factors to be controlled to ensure the validity of the test and calibration results.</p> <p><i>NOTE 1: Sampling is a defined procedure whereby a part of a substance, material or product is taken to provide for testing or calibration of a representative sample of the whole. Sampling may also be required by the appropriate specification for which the substance, material or product is to be tested or calibrated. In certain cases (e.g. forensic analysis), the sample may not be representative but is determined by availability.</i></p> <p><i>NOTE 2: Sampling procedures should describe the allocation, sampling plan, withdrawal and preparation of a sample or samples from a substance, material or product to yield the required information.</i></p> | <p>In general there is no sampling in the sense of ISO/IEC 17025 in type evaluation process in legal metrology.</p> <p>Nevertheless, OIML Recommendations may require a selection of samples amongst a family of measuring instruments.</p> <p>Such a selection is done under the responsibility of the Issuing Authority.</p> <p>If sampling is requested for legal metrology control, it is under the responsibility of the Issuing Authority.</p> |
| <p>5.7.2 Where the customer requires deviations, additions or exclusions from the documented sampling procedure, these shall be recorded in detail with the appropriate sampling data and shall be included in all documents containing test and/or calibration results, and shall be communicated to the appropriate personnel.</p> | <p>Not applicable (see 5.7.1)</p> |
| <p>5.7.3 The laboratory shall have procedures for recording relevant data and operations relating to sampling that forms part of the testing or calibration that is undertaken. These records shall include the sampling procedure used, the identification of the sampler, environmental conditions (if relevant) and diagrams or other equivalent means to identify the sampling location as necessary and, if appropriate, the statistics the sampling procedures are based upon.</p> | |

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| <p>5.8 Handling of test and calibration items</p> | |
| <p>5.8.1 The laboratory shall have procedures for the transportation, receipt, handling, protection, storage, retention and/or disposal, of test and/or calibration items, including all provisions necessary to protect the integrity of the test or calibration item, and to protect the interests of the laboratory and the customer.</p> | <p>Provisions shall be made to ensure that the sample to be examined and/or tested is the one validated by the Issuing Authority. Such provisions may consist in receiving the sample from the Issuing Authority or in receiving it in sealed packaging. In this case, the Testing Laboratory shall register the seal position when the packaging is received.</p> |
| <p>5.8.2 The laboratory shall have a system for identifying test and/or calibration items. The identification shall be retained throughout the life of the item in the laboratory. The system shall be designed and operated so as to ensure that items cannot be confused physically or when referred to in records or other documents. The system shall, if appropriate, accommodate a sub-division of groups of items and the transfer of items within and from the laboratory.</p> | |
| <p>5.8.3 Upon receipt of the test or calibration item, abnormalities or departures from normal or specified conditions, as described in the test or calibration method, shall be recorded. When there is doubt as to the suitability of an item for test or calibration, or when an item does not conform to the description provided, or the test or calibration required is not specified in sufficient details, the laboratory shall consult the customer for further instruction before proceeding and shall record the discussion.</p> | <p>The procedures of the Testing Laboratory shall indicate that the Issuing Authority is consulted in such a case.</p> |
| <p>5.8.4 The laboratory shall have procedures and appropriate facilities for avoiding deterioration, loss or damage to the test or calibration item during storage, handling and preparation. Handling instructions provided with the item shall be followed. When items have to be stored or conditioned under specified environmental conditions, these conditions shall be maintained, monitored and recorded. Where a test or calibration item or portion of an item is to be held secure, the laboratory shall have arrangements for storage and security that protect the condition and integrity of the secured items or portions concerned.</p> <p><i>NOTE 1: Where test items are to be returned into service after testing, special care is required to ensure that they are not damaged or injured during the handling, testing or storing/waiting processes.</i></p> <p><i>NOTE 2: A sampling procedure and information on storage and transport of samples, including information on sampling factors influencing the test or calibration result, should be provided to those responsible for taking and transporting the samples.</i></p> <p><i>NOTE 3: Reasons for keeping a test or calibration item secure can be for reasons of record, safety or value, or to enable complementary tests and/or calibrations to be performed later.</i></p> | <p>Provisions for transportation, conditioning, handling, and installation for testing shall be determined with the agreement of the Issuing Authority.</p> |

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| <p>5.9 Assuring the quality of test and calibration results</p> | |
| <p>5.9.1 The laboratory shall have quality control procedures for monitoring the validity of tests and calibrations undertaken. The resulting data shall be recorded in such a way that trends are detectable and, where practicable, statistical techniques shall be applied to the reviewing of the results. This monitoring shall be planned and reviewed and may include, but not be limited to, the following:</p> <ul style="list-style-type: none"> a) regular use of certified reference materials and/or internal quality control using secondary reference materials; b) participation in interlaboratory comparison or proficiency-testing-programmes; c) replicate tests or calibrations using the same or different methods; d) retesting or recalibration of retained items; e) correlation of results for different characteristics of an item. <p><i>NOTE: The selected methods should be appropriate for the type and volume of the work undertaken.</i></p> | <p>Such procedures shall include participation in interlaboratory comparisons organized by the BIML, if necessary.</p> |
| <p>5.9.2 Quality control data shall be analysed and, where they are found to be outside pre-defined criteria, planned action shall be taken to correct the problem and to prevent incorrect results from being reported.</p> | |
| <p>5.10 Reporting the results</p> | |
| <p>5.10.1 General</p> <p>The results of each test, calibration, or series of tests or calibrations carried out by the laboratory shall be reported accurately, clearly, unambiguously and objectively, and in accordance with any specific instructions in the test or calibration methods.</p> <p>The results shall be reported, usually in a test report or a calibration certificate (see Note 1), and shall include all the information requested by the customer and necessary for the interpretation of the test or calibration results and all information required by the method used. This information is normally that required by clause 5.10.2, and 5.10.3 or 5.10.4.</p> <p>In the case of tests or calibrations performed for internal customers, or in the case of a written agreement with the customer, the results may be reported in a simplified way. Any information listed in clause 5.10.2 to 5.10.4 which is not reported to the customer shall be readily available in the laboratory which carried out the tests and/or calibrations.</p> <p><i>NOTE 1: Test reports and calibration certificates are sometimes called test certificates and calibration report, respectively.</i></p> | <p>Each Testing Laboratory issues a Partial Test Report or an OIML Partial Test Report and/or an Additional Test Report which corresponds to tests and examinations it performed.</p> <p>Relevant parts of the Test Report Format given in the applicable OIML Recommendation shall be used by each Testing Laboratory for issuing the OIML Partial Test Report.</p> <p>The results of examinations in the OIML Partial Test Report shall be provided using the applicable checklists in the relevant OIML Recommendation.</p> <p>The Test Report (Partial Test Report, OIML Partial Test Report, Additional Test Report) shall indicate:</p> <ul style="list-style-type: none"> - whether the tests carried out were split up between two or more samples, - whether adjustments and/or modifications were performed during tests, - the reason why some tests were not performed, in particular in the case of a complementary Type Approval process. <p>In the event of the MAA implementation, the Test Report Format specified in the applicable procedures validated by the CPR shall be used to issue the Additional Test Report(s).</p> |

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| <p><i>NOTE 2: The test reports or calibrations certificates may be issued as hard copy or by electronic data transfer provided that the requirements of this Standard are met.</i></p> | <p>Third paragraph: Not applicable.</p> |
| <p>5.10.2 Test reports and calibration certificates Each test report or calibration certificate shall include at least the following information, unless the laboratory has valid reasons for not doing so:</p> <ul style="list-style-type: none"> a) a title, (e.g. "Test Report" or 'Calibration Certificate'); b) the name and address of the laboratory, and the location where the tests and/or calibrations were carried out, if different from the address of the laboratory; c) unique identification of the test report or calibration certificate (such as the serial number), and on each page an identification in order to ensure that the page is recognized as a part of the test report or calibration certificate, and a clear identification of the end of the test report or calibration certificate; d) the name and address of the customer; e) identification of the method used; f) a description of, the condition of, and unambiguous identification of the item(s) tested or calibrated; g) the date of receipt of the test or calibration item(s) where this is critical to the validity and application of the results, and the date(s) of performance of the test or calibration; h) reference to the sampling plan and procedures used by the laboratory or other bodies where these are relevant to the validity or application of the results; i) the test or calibration results with, where appropriate, the units of measurement; j) the name(s), function(s) and signature(s) or equivalent identification of person(s) authorizing the test report or calibration certificate; k) where relevant, a statement to the effect that the results relate only to the items tested or calibrated. <p><i>NOTE 1: Hard copies of test reports and calibration certificates should also include the page number and total number of pages.</i></p> <p><i>NOTE 2: It is recommended that laboratories include a statement specifying that the test report or calibration certificate shall not be reproduced except in full, without written approval of the laboratory.</i></p> | <ul style="list-style-type: none"> f) Includes where tests were split up between several samples. h) Testing Laboratories shall indicate the verifications performed to ensure that the tested and/or examined samples are those validated by the Issuing Authority. |

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| <p>5.10.3 Test reports</p> | |
| <p>5.10.3.1 In addition to the requirements listed in 5.10.2 test reports shall, where necessary for the interpretation of the test results, include the following:</p> <ul style="list-style-type: none"> a) deviations from, additions to or exclusions from the test method, and information on specific test conditions, such as environmental conditions; b) where relevant, a statement of compliance/ non compliance with requirements and/or specifications; c) where applicable, a statement on the estimated uncertainty of measurement; information on uncertainty is needed in test reports when it is relevant to the validity or application of the test results, when a customer's instruction so requires, or when the uncertainty affects compliance to a specification limit; d) where appropriate and needed, opinions and interpretations (see 5. 10. 5); e) additional information which may be required by specific methods, customers or group of customers. | <ul style="list-style-type: none"> b) The Test Report Format of the relevant OIML Recommendation specifies the way in which the interpretations shall be indicated (pass/fail). c) The information may be provided in the form of a ratio between the expanded uncertainty and the maximum permissible error. The coverage factor shall always be indicated. e) For example, information related to the environmental conditions under which the tests were performed outdoors (see 5.3.2). |
| <p>5.10.3.2 In addition to the requirements listed in 5.10.2 and 5.10.3.1, test reports containing the results of sampling shall include the following, where necessary for the interpretation of test results:</p> <ul style="list-style-type: none"> a) the date of sampling; b) unambiguous identification of the substance, material or product sampled (including the name of the manufacturer, the model or type of designation and serial numbers as appropriate); c) the location of sampling, including any diagrams, sketches or photographs; d) a reference of the sampling plan and procedures used; e) details of any environmental conditions during sampling that may affect the interpretation of the test results; f) any standard or other specification for the sampling method or procedure, and deviations, additions to or exclusions from the specification concerned. | |
| <p>5.10.4 Calibration certificates</p> | |
| <p>5.10.4.1 In addition to the requirements listed in 5.10.2 calibration certificates shall include the following, where necessary for the interpretation of the calibration results:</p> <ul style="list-style-type: none"> a) the conditions (e.g. environmental) under which the calibrations were made that have an influence on the measurement results; b) the uncertainty of measurement and/or a statement of compliance with an identified metrological specification or clauses thereof; c) evidence that the measurements are traceable (see note 2 in 5.6.2.1.1). | <p>Not applicable.</p> |

| <p style="text-align: center;">ISO 17025 General requirements for the competence of testing and calibration laboratories to testing laboratory activities</p> | <p style="text-align: center;">Specific requirements and comments related to Testing Laboratories involved in legal metrology testing</p> |
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| <p>5.10.4.2 The calibration certificate shall relate only to quantities and the results of functional tests. If a statement of compliance with a specification is made, this shall identify which clauses of the specification are met or not met.</p> <p>When a statement of compliance with a specification is made omitting the measurement results and associated uncertainties, the laboratory shall record those results and maintain them for possible future reference.</p> <p>When statements of compliance are made, the uncertainty of measurement shall be taken into account.</p> | <p>Not applicable.</p> |
| <p>5.10.4.3 When an instrument for calibration has been adjusted or repaired, the calibration results before and after adjustment or repair, if available, shall be reported.</p> | <p>Not applicable.</p> |
| <p>5.10.4.4 A calibration certificate (or calibration label) shall not contain any recommendation on the calibration interval except where this has been agreed with the customer. This requirement may be superseded by legal regulations.</p> | <p>Not applicable.</p> |
| <p>5.10.5 Opinions and interpretations</p> <p>When opinions and interpretations are included, the laboratory shall document the basis upon which the opinions and interpretations have been made. Opinions and interpretations shall be clearly marked as such in a test report.</p> <p><i>NOTE 1: Opinions and interpretations should not be confused with inspections and product certifications as intended in the ISO/IEC 17020 and ISO/IEC Guide 65.</i></p> <p><i>NOTE 2: Opinions and interpretations included in a test report may comprise, but not be limited to, the following:</i></p> <ul style="list-style-type: none"> - an opinion on the statement of compliance/non compliance of the results with requirements; - fulfilment of contractual requirements; - recommendations on how to use the results; - guidance to be used for improvements. <p><i>NOTE 3: In many cases it might be appropriate to communicate the opinions and interpretations by direct dialogue with the customer. Such dialogue should be written down.</i></p> | <p>See 5.10.3.1 d)</p> <p>Opinions and interpretations related to the conformance of the instrument with the relevant OIML Recommendation are not allowed in Test Reports (Partial Test Report, OIML Partial Test Report, Additional Test Report).</p> <p>The Issuing Authority is solely responsible for drawing conclusions on the conformance of the instrument with the relevant requirements (e.g. OIML Recommendation)</p> |
| <p>5.10.6 Testing and calibration results obtained from sub-contractors</p> <p>When the test report contains results of tests performed by sub-contractors, these results shall be clearly identified. The subcontractor shall report the results in writing or electronically.</p> | <p>In the event of tests and/or examinations performed by several Testing Laboratories, each Testing Laboratory shall establish a Test Report (Partial Test Report, OIML partial Test Report, Additional Test Report) . Each Test Report is then a part of the Evaluation Report or OIML Evaluation Report and, if applicable, Complete Evaluation Report.</p> |

| <p style="text-align: center;">ISO 17025 General requirements for the competence of testing and calibration laboratories to testing laboratory activities</p> | <p style="text-align: center;">Specific requirements and comments related to Testing Laboratories involved in legal metrology testing</p> |
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| <p>When a calibration has been subcontracted, the laboratory performing the work shall issue the calibration certificate to the contracting laboratory.</p> | |
| <p>5.10.7 Electronic transmission of results</p> <p>In the case of transmission of test or calibration results by telephone, telex, facsimile or other electronic or electromagnetic means, the requirements of this International Standard shall be met (see also 5.4.7).</p> | |
| <p>5.10.8 Format of reports and certificates</p> <p>The format shall be designed to accommodate each type of test or calibration carried out and to minimize the possibility of misunderstanding or misuse.</p> <p><i>NOTE 1: Attention should be given to the lay-out of the test report or calibration certificate, especially with regard to the presentation of the test or calibration data and ease of assimilation by the reader.</i></p> <p><i>NOTE 2: The headings should be standardized as far as possible.</i></p> | <p>See 5.10.1.</p> |
| <p>5.10.9 Amendments to test reports and calibration certificates</p> <p>Material amendments to a test report or calibration certificate after issue shall be made only in the form of a further document, or data transfer, which includes the statement:</p> <p>“Supplement to Test Report [or Calibration Certificate], serial number... [or as otherwise identified]”, or an equivalent form of wording.</p> <p>Such amendments shall meet all the requirements of this International Standard.</p> <p>When it is necessary to issue a complete new test report or calibration certificate, this shall be uniquely identified and shall contain a reference to the original that it replaces.</p> | |