

[Organization's Logo]

[Organization's Name]

**Commented [170251]:** All fields in this document marked by square brackets [ ] must be filled in.

## CALIBRATION REPORT AND CERTIFICATE REQUIREMENTS PROCEDURE

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Date of version:	
Signature:	

**Commented [170252]:** Adapt to the existing practice in organization.

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**Commented [17A3]:** This is only necessary if document is in paper form; otherwise, this table should be deleted.

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## Change History

Date	Version	Created by	Description of change
	0.1	17025Academy	Basic document outline

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## 1. Purpose, Scope and Users

The purpose of this procedure is to outline the requirements for writing calibration reports and certificates that meet the requirements of the international standard.

This procedure applies to all calibration reports and certificates issued for both external and internal use.

Users of this procedure are laboratory personnel who provide final calibration reports to customers.

## 2. References

- ISO/IEC 17025:2017; Clause 7.8.2, 7.8.4, 7.8.6, 7.8.7
- Quality Manual
- Sampling Procedure
- Document and Record Control Procedure
- JCGM 200: 2012 International vocabulary of metrology

## 3. Calibration Report Procedure

### 3.1. Calibration Reports and Certificates

[Job title] must ensure that all calibration reports and certificates are provided to the customer

in a simplified way to meet the customer's needs and requirements. Information required by the

- Calibration report or certificate title.
- The name and address of the laboratory.
- [Redacted]
- [Redacted]
- [Redacted]
- Identification of the method used.

**Commented [17A4]:** You will find this document in the ISO 17025 Toolkit folder "03\_Quality\_Manual".

**Commented [17A5]:** You will find this document in the ISO 17025 Toolkit folder "12\_Sampling".

**Commented [17A6]:** You will find this document in the ISO 17025 Toolkit folder "00\_Document\_and\_Record\_Control".

**Commented [170257]:** Quality manager or person designated by Quality manager.

**Commented [170258]:** Customer can decide what sort and presentation of results wants to have on a report.

**Commented [170259]:** The form of calibration report

**Commented [1702510]:** Laboratory or Quality manager.

**Commented [1702511]:** In case of deviations or exclusions from the method or external providers etc.

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- A description, unambiguous identification and, when necessary, the condition of the item.
- The date of receipt of the substance or the date of sampling, when this is related to the validity and application of the results.
- The nature of performance of the substance, material or product.
- The date of issue of the report or certificate.
- Reference to the sampling and measurement methods used, the laboratory or other bodies when these are relevant to the validity or application of the results.
- A statement to the effect that the results relate only to the items calibrated.
- The calibration results with units of measurement, when applicable.
- Reference to, location, or extension from the report.
- Identification of the person authoring the report.
- Any identification other results or their related results.

It is the responsibility of the [job title] to ensure all calibration reports and certificates meet all requirements and the responsibility of all laboratory personnel to gather and record calibration

**Commented [1702512]:** Laboratory or Quality manager.

identified. In addition, a disclaimer must be put on the report or certificate when information is supplied by the customer and can affect the validity of the calibration results.

**Commented [1702513]:** Laboratory or Quality manager.

When [organization name] has not been responsible for the sampling stage (e.g. the sample was

When [organization name] is not responsible for the sampling stage, the [job title] issuing results of sampling must address the following information:

**Commented [1702514]:** In case when necessary for the interpretation of the calibration results according to sampling procedure.

- The date of sampling.
- Unambiguous identification of the substance, material or product sampled including the name of the manufacturer, the trade or type of preparation and any number or identifier.
- The location of sampling, including any relevant details or photographs.
- A reference to the sampling and measurement methods used, the laboratory or other bodies when these are relevant to the validity or application of the results.
- Information required to evaluate measurement uncertainty for subsequent testing.

[organization name]

- The measurement uncertainty of the result presented in the same unit as that of the measurand or in a term relative to the measurand (e.g. percent).
- The measurement uncertainty is expressed in terms of the measurand and not in terms of the reference value.
- A statement identifying the measurement uncertainty is provided for each of the measurands.
- The measurement uncertainty is expressed in terms of the measurand.

A calibration certificate or calibration must will not contain any recommendation on the calibration interval except where this has been agreed with the customer.

### 3.2. Statements of conformity

When a statement of conformity to a specification or standard for calibration is provided by

[redacted]

Commented [1702515]: Laboratory or Quality manager.

[Job title] reports on the statement of conformity such that the statement clearly identifies:

Commented [1702516]: Laboratory or Quality manager.

- To which results the statement applies
- The measurement uncertainty associated with the statement
- The measurand to which the statement applies

When reported information associated with a calibration includes a statement of conformity with a specification, omitting the measurement results and associated uncertainties, the reported information

[redacted]

### 3.3. Opinions and Interpretations

The opinions and interpretations expressed in calibration reports and certificates must be based on the

[redacted]

[Job title] documents the information upon which the opinions and interpretations have been based.

Commented [1702517]: Laboratory or Quality manager.

[Job title] communicates opinions and interpretations with the customer and keep a record of the

Commented [1702518]: An example would be when an

[redacted]

### 3.4. Amendments to Reports

When a report is amended, the change should be clearly identified and, where appropriate, the reason for the change should be included in the report.

Commented [1702519]: Quality manager or Laboratory manager.

Commented [1702520]: Other documents, which must all

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Amendments to a report after issue must only be made by [job title] in the form of a further document or data transfer that includes the statement:

Amendment to Calibration Report and Certificate [1702521] is an amendment form of [1702521]

This amendment is not valid and has been removed from use.

original that is no longer valid and has been removed from use.

**Commented [1702521]:** Laboratory or Quality manager.

**Commented [1702522]:** Use in case of other principle of identification of the original Calibration Report.

#### 4. Calibration Records

All calibration reports and copies of certificates become official technical records, both hardcopies and electronic copies and are kept on file indefinitely to support regulatory and customer requirements.

Records may be stored electronically. If the record is in electronic form, write the name of the folder on Laboratory Manager's computer.

#### 5. Managing Records Kept on the Basis of this Document

Document or record name	Identification	Storage		Responsibility
		Retention time	Location	
Calibration Report	Customer files	Permanent	[office of [job title]]	[job title]
Calibration Records	Customer files	Permanent	[office of [job title]]	[job title]
Copies of issued certificates	Customer files	Permanent	[office of [job title]]	[job title]

**Commented [1702523]:** Calibration reports and certificates are issued on formal organization letterhead and include all the requirements of the standard with reference to the accreditation registrar and their logo.

Records may be stored electronically. If the record is in electronic form, write the name of the folder on Laboratory Manager's computer.

**Commented [1702524]:** These records are kept by the customer and the retention times are set by them.

Electronic copies kept by the calibration may be kept indefinitely.

**Commented [1702525]:** If the record is in electronic form, write the name of the folder on [job title]'s computer.

**Commented [1702526]:** The content of calibration report template is within the domain of the organization.

**Commented [1702528]:** Assigned to the person who is responsible for document and records control in the laboratory.

**Commented [1702527]:** E.g. Calibration Laboratory Office for copies kept by the calibration laboratory.

**Commented [1702529]:** Each laboratory organization makes own rules and requirements about writing of calibration records, compulsory by each testing method or calibration.

**Commented [1702530]:** The content of certificate is within the domain of the organization.