

ISoOR-AB

Accreditation Process Procedure

1. Purpose

This procedure defines the process by which ISoOR-AB evaluates and accredits Conformity Assessment Bodies (CABs), ensuring compliance with applicable standards and accreditation requirements.

2. Scope

This procedure applies to all accreditation activities conducted by ISoOR-AB, including initial accreditation, surveillance, and re-accreditation.

3. Process Overview

The accreditation process consists of the following stages:

1. Application
 2. Application Review
 3. Assessment Planning
 4. Document Review
 5. On-site / Remote Assessment
 6. Nonconformity Management
 7. Accreditation Decision
 8. Surveillance and Re-accreditation
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4. Application

- CAB submits a formal application including:

- Scope of accreditation
 - Quality management documentation
 - Organizational structure
 - ISOOR-AB acknowledges receipt and assigns a case manager
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5. Application Review

- Completeness and eligibility are evaluated
 - Conflicts of interest are checked
 - Decision to proceed or reject application
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6. Assessment Planning

- Assessment team is appointed
 - Competence of assessors is verified
 - Assessment plan is communicated to the CAB
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7. Document Review

- Evaluation of:
 - Quality system
 - Procedures
 - Technical competence
 - Identification of potential gaps prior to assessment
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8. Assessment

- Conducted on-site or remotely
- Includes:
 - Interviews
 - Witnessing activities
 - Record review

9. Nonconformity Management

- Findings are classified (e.g., major, minor)
 - CAB must submit corrective actions within defined timelines
 - Verification of corrective actions by ISO-AB
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10. Accreditation Decision

- Decision made by independent decision-makers
 - Assessment team shall not make the final decision
 - Possible outcomes:
 - Accreditation granted
 - Accreditation granted with conditions
 - Accreditation denied
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11. Surveillance

- Periodic monitoring (e.g., annually)
 - Includes document review and/or reassessment
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12. Re-accreditation

- Full reassessment conducted at defined cycle (e.g., every 3–5 years)
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13. Records

- All stages shall be documented and retained

