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***Publication  
Reference***

**EA-INF/01: 2017**

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# **List of EA Publications And International Documents**

## ***PURPOSE***

This publication gives the list of EA documents to be used either on a mandatory or informative basis, according to the classification given in EA-1/14 “*Procedure for Development and Approval of EA Documents and the Adoption of ILAC/IAF Documents*”. It contains also the list of International documents approved by EA as applicable EA documents.

*Authorship*

This document has been prepared by the EA Secretariat.

*Official language*

The publication may be translated into other languages as required. The English language version remains the definitive version.

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*Further information*

For further information about this publication, please contact the EA Secretariat at: [secretariat@european-accreditation.org](mailto:secretariat@european-accreditation.org)

Please check our website for up-to-date information at <http://www.european-accreditation.org>

**Category:** Information and Promotional Documents

**Date of implementation:** Immediate

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**CLASSIFICATION OF DOCUMENTS**

At its meeting in May 2009, the EA General Assembly decided to change the classification of the documents. EA Publications are now classified in 6 categories. Within one category, documents can have different status as shown below:

Categories	Status	Numbering
<b>Secretariat Management System and related Documents</b>	Not Applicable	EA-0/XX
<b>Information and Promotional documents</b>	Not Applicable	EA-INF/XX
<b>EA Governance and Policy documents</b>	Not Applicable	<b>A:</b> documents relevant for EA as an Association :EA-1/XX <b>A</b> <b>AB:</b> documents relevant for Members: EA-1/XX <b>AB</b>
<b>Peer Evaluation Process documents, including policies and procedures</b>	Not Applicable	EA-2/02
<b>Members' Procedural documents</b>	<ul style="list-style-type: none"> <li>• <b>Mandatory</b></li> <li>or</li> <li>• <b>Guidance</b></li> <li>or</li> <li>• <b>/INF</b>ormative documents</li> </ul>	EA-2/XX <b>M</b> and EA-3/XX <b>M</b> or EA-2/XX <b>G</b> and EA-3/XX <b>G</b> EA-2/XX <b>INF</b> and EA-3/XX <b>INF</b>
<b>Application documents and Technical/Advisory for Conformity Assessment Bodies</b>	<ul style="list-style-type: none"> <li>• <b>Mandatory</b></li> <li>or</li> <li>• <b>Guidance</b></li> <li>or</li> <li>• <b>INF</b>ormative</li> <li>or</li> <li>• <b>Technical / Advisory</b> documents</li> </ul>	EA-4/XX <b>M</b> and EA-5/XX <b>M</b> EA-6/XX <b>M</b> and EA-7/XX <b>M</b> EA-4/XX <b>G</b> and EA-5/XX <b>G</b> EA-6/XX <b>G</b> and EA-7/XX <b>G</b> EA-4/XX <b>INF</b> and EA-5/XX <b>INF</b> EA-6/XX <b>INF</b> and EA-7/XX <b>INF</b> EA-4/XX <b>TA</b> and EA-5/XX <b>TA</b> EA-6/XX <b>TA</b> and EA-7/XX <b>TA</b> and other Organisations.

**EA Mandatory Documents** are required to be used by accreditation bodies when accrediting conformity assessment bodies to ensure that both operate their activities in a consistent and equivalent manner. EA Mandatory Documents do not establish, interpret, subtract from or add to the requirements of any ISO/IEC/EN Guides and Standards or other technical specifications and normative documents, but assure consistent application of those documents.

For further information on the categories of EA documents and its classification into Mandatory, Guidance, Informative and Technical/Advisory documents, see EA-1/14 Procedure for Development and Approval of EA Documents and the Adoption of ILAC/IAF Documents.

## **1 DOCUMENT CONTROL**

As owners of the EA publications, the EA Committees are responsible for the drafting, control and revision of their documents.

The same applies for the documents owned by the EA Secretariat.

The EA Secretariat is responsible for the preparation of the documents for publication on the EA web site according to EA-0/06 Format and Layout of EA Documents and EA-1/14 - Procedure for Development and Approval of EA Documents and the Adoption of ILAC/IAF Documents.

Periodically, at least every 5 years, the Committees shall review the publications they are responsible for.

For any question relating to a publication, please contact the Committee in charge or the EA Secretariat.

## 2 EA PUBLICATIONS

### 2.1 SECRETARIAT MANAGEMENT SYSTEM AND RELATED DOCUMENTS

Reference	Title	Date of publication	Date end transition (blank if not applicable)
EA-0/00 (rev.01)	EA Management System	May 2015	
EA-0/02 (rev.00)	Procedure for Document Control	Sept 2012	
EA-0/06 (rev.02)	Format and Layout of EA Documents	May 2009	
EA-0/07 (rev.01)	EA Procedure for Application for EA Membership and Application Form	May 2017	
EA-0/08 (rev.00)	Procedure for management reviews	July 2017	
EA-0/09 (rev.01)	Procedure for internal audits	Oct 2015	
EA-0/10 (rev.01)	Procedure for EA Meetings	Sept 2012	
EA-0/11 (rev.00)	Specific Provisions for the Implementation of the Peer Evaluation of National Accreditation Bodies according to EN ISO 14065 and Commission Regulation (EU) No 600/2012	March 2014	
EA-0/12 (rev.00)	Guideline for implementation of EA Projects	June 2015	

### 2.2 INFORMATION AND PROMOTIONAL DOCUMENTS

*Information and promotional publications such as EA brochures, annual reports, activity reports and other information on EA and EA activities. These documents are prepared and published by the Secretariat or EA Committees/Councils.*

Reference	Title	Date of publication	Date end transition (blank if not applicable)
EA-INF/01	List of EA Publications (this publication)	07 August 2017	
EA-INF/02 (rev.21)	Contact Persons of EA Members, Associate Members, Recognized Stakeholders and Observers	August 2017	

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EA-INF/03 (rev.61)	EA Multilateral and Bilateral Agreements, and Signatory Lists	October 2017
EA-INF/04 (rev.04)	Statement on acceptance and recognition of activities under the EA MLA previously EA-1/16	May 2016
EA-INF/05 (rev.01)	Press kit <i>Document temporarily withdrawn</i>	Temporarily withdrawn
EA-INF/06 (rev.01)	A Guide to EA <i>Document under review</i>	Document under revision
EA-INF/07 (rev.01)	Accreditation Body – Communication with National Regulators - Best Practice Guide	June 2011
EA-INF/08 (rev.01)	A Briefing for Government and Regulators	2013
EA-INF/09 (rev.00)	EA Guide on the Publication of Data on Accredited Verifiers	Jan 2014
EA-INF/10 (rev.00)	Guide on the Content of the Accreditation Certificate for Verifiers Accredited for EN ISO 1465 and the Commission Regulation (EU) No 600/2012	Jan 2014
EA-INF/11 (rev.00)	Scope of the EA MLA and the application of an EA MLA signatory to join the IAF MLA for a specific main scope/sub-scope and/or the ILAC MRA for a specific activity	Apr 2014
EA-INF/12 (rev00)	Benefits and importance of the participation in EA highlighted PT schemes	Withdrawn May 2017
EA-INF/13 (rev00)	The Assessment and Accreditation of Opinions and Interpretations using ISO/IEC 17025:2005	Sept 2015
EA-INF/14 (rev00)	Communication with the European Commission and its DG's	Oct 2015
EA-INF/15 (rev00)	Joint EA-EDQM Communication regarding Cooperation when carrying out (joint) audits/assessments in Official Medicines Control Laboratories	Apr 2017

## 2.3 GOVERNANCE AND POLICY DOCUMENTS

Documents are relating to the operation of EA as an association and a legal entity.

The following letters are added to the document number:

**A:** documents relevant for EA as an Association e.g. EA-1/XX **A**

**AB:** documents relevant for EA Members e.g. EA-1/XX **AB**

Reference	Title	Date of publication	Date end transition (blank if not applicable)
EA-1/06 A+AB (rev.07)	EA Multilateral Agreement Criteria for signing Policy and procedure for development	Oct 2015 (Re-published)	
EA-1/13 A (rev.01)	EA's Relationship with Accreditation Bodies of Countries Not Being Members of the EU or EFTA	May 2016	
EA-1/14 A (rev.01)	Procedure for Development and Approval of EA Documents and the Adoption of ILAC/IAF Documents	May 2017	
EA-1/15 A (rev.01)	EA Policy for Relations with Stakeholders	Aug 2015	
EA-1/17 A (rev.06)	EA Rules of Procedure	June 2017	
EA-1/17 S1 A+AB (rev.03)	Supplement 1 to EA-1/17, Criteria for Membership	June 2014	
EA-1/17 S3 A (rev.04)	Supplement 3 to EA-1/17, Complaint and Appeal Procedures	May 2017	
EA-1/17 S4 A (rev.01)	Supplement 4 to EA-1/17, Proxy Procedure	May 2017	
EA-1/17 S5 A+AB (rev.03)	Supplement 5 to EA-1/17, Levying of Membership Fees	November 2016	
EA-1/19 A+AB (rev.02)	Rules for Use of EA logo and Graphic Specification	April 2016	
EA-1/20 A (rev.03)	EA Procedure for the Control of Expenditures and Preparation of Budgets (previously EA-0/05)	May 2016	



EA-1/20 S1 A+AB (rev.04)	EA Supplement 1 to EA-1/20. Terms and Conditions for financial compensation from the operating/action grant to an EA Member Accreditation Body	May 2016
EA-1/21 A+AB (rev.00)	EA Internal Procedure for Liaison Activities ( <i>previously EA-0/04</i> )	July 2008
EA-1/22 A+AB (rev.03)	EA Procedure and Criteria for the Evaluation of Conformity Assessment Schemes (Previously EA-2/11)	Nov 2016

## 2.4 PEER EVALUATION PROCESS DOCUMENTS

*Policy and procedural documents for the operation and management of the EA peer evaluation system, training of evaluators and liaison with the Arrangement Committees of ILAC and IAF.*

Reference	Title	Date of publication	Date end transition (blank if not applicable)
EA-2/02 M (rev.07)	EA Procedure for the evaluation of a National Accreditation Body	May 2016	
EA-2/02 S1 (rev.00)	EA Supplement 1 to EA-2/02. Selection, Training and Monitoring of Evaluators	Jan 2016	

## 2.5 MEMBERS' PROCEDURAL DOCUMENTS

*Procedures, requirements etc. applying to EA members including application documents for accreditation bodies.*

Documents in this section can be either mandatory (**M**) or guidance (**G**) or informative (**INF**) document

Reference	Classification	Title	Date of publication	Date end transition (blank if not applicable)
EA-2/13 M (rev.01)	Mandatory	EA Cross Border Accreditation Policy and Procedure for Cross Border Cooperation between EA Members	Oct 2012	
EA-2/13 S1 M (rev.00)	Mandatory	Supplement 1 to EA-2/13, Interpretation of Terminology used in clause 5.1 of EA-2/13	Aug 2013	
EA-2/14 M (rev.00)	Mandatory	Procedure for Regional Calibration ILCs in Support of the EA MLA	Withdrawn May 2017	
EA-2/15 M	Mandatory	EA Requirements for the Accreditation of Flexible Scopes	July 2008	

(rev.00)				
EA-2/17 M (rev.03)	Mandatory	EA Document on Accreditation for Notification purposes	November 2016	
EA-2/18 INF (rev.00)	Informative	Guidelines for Accreditation Bodies on the content of the scopes of accreditation for proficiency testing providers	Oct 2015	
EA-3/01 M (rev.03)	Mandatory	EA conditions for the use of accreditation symbols, text reference to accreditation and MLA signatory status	Dec 2012	
EA-3/04 G (rev.01)	Guidance	Use of Proficiency Testing as a Tool for Accreditation in Testing ( <i>with EUROLAB and EURACHEM</i> ). Reinstated according to EA Resolution 2012(29)24	Aug 2001	
EA-3/11 M (rev.00)	Mandatory	Food Safety Management Systems – Scope of Accreditation	May 2009	
EA-3/12 M (rev.00)	Mandatory	EA Policy for Accreditation of Organic Production Certification	June 2013	
EA-3/13 M (rev.00)	Mandatory	EA Document on the Application of ISO/IEC 17021-1 for the Certification of Occupational Health and Safety Management Systems	June 2016	
EA-5/02 INF (rev.02)	Informative	EA Guidance on the Application of ISO/IEC 17020 in Periodic Inspection of the Roadworthiness of Motor Vehicles and their Trailers	Dec 2015	

## 2.6 APPLICATION DOCUMENTS AND TECHNICAL/ADVISORY DOCUMENTS FOR CONFORMITY ASSESSMENTS BODIES (CABs)

*Documents used by Conformity Assessment Bodies providing technical or scientific guidance for the application of standards.*

The documents of this category can be either:

Mandatory (**M**) or Guidance (**G**) or Informative (**INF**) or Technical/Advisory (**TA**)

### 2.6.1 Laboratories - ISO/IEC 17025 / ISO 15189

Reference	Classification	Title	Date of publication	Date end transition (blank if not applicable)
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EA-4/02 M (rev.01)	Mandatory	Expressions of the Uncertainty of Measurements in Calibration (including supplement 1 to EA-4/02) ( <i>previously EAL- R2</i> )	Sept 2013
EA-4/09 G (rev.02)	Guidance	Accreditation for Sensory Testing Laboratories ( <i>previously EAL-G16</i> )	Feb 2017
EA-4/14 INF (rev.00)	Informative	Selection and Use of Reference Materials	Feb 2003
EA-4/15 G (rev.01)	Guidance	Accreditation for Non-Destructive Testing	May 2015
EA-4/16 G (rev.00)	Guidance	EA Guidelines on the Expression of Uncertainty in Quantitative testing	Dec 2003
EA-4/17 M (rev.00)	Mandatory	EA Position Paper on the description of scopes of accreditation of medical laboratories	Dec 2008
EA-4/18 INF (rev.00)	Informative	Guidance on the level and frequency of proficiency testing participation	June 2010
EA-4/20 G	Guidance	Guidance for the Assessment of Laboratories against EN ISO 15189 and 22870	May 2015
EURACHEM	Technical / Advisory	Guidance on Accreditation of Microbiological Laboratories (2013)	2013
CEC TA	Technical / Advisory	ISO/IEC 17025 interpretation document for CEC test methods Coordination European Council: Ian Brown	Aug 2006
CITAC / EURACHEM TA	Technical / Advisory	Guide to Quality in analytical Chemistry	July 2005
EDQM/OMCL TA	Technical / Advisory	Validation of analytical procedures	Feb 2014
EDQM/OMCL TA	Technical / Advisory	Scope of accreditation of official medicines laboratories	Dec 2007
EDQM/OMCL TA	Technical / Advisory	Uncertainty of measurement	Dec 2007
EDQM/OMCL TA	Technical / Advisory	Standard – Aide – Mémoire for the Mutual Joint Audit of OMCLs	Feb 2011

EDQM/OMCL TA	Technical / Advisory	Qualification of Equipment (core document)	July 2011
EDQM/OMCL TA	Technical / Advisory	Annex 1: Qualification of HPLC equipment	July 2011
EDQM/OMCL TA	Technical / Advisory	Annex 2: Qualification of GC equipment	Oct 2006
EDQM/OMCL TA	Technical / Advisory	Annex 3: Qualification of UV – visible spectrophotometers	Dec 2007
EDQM/OMCL TA	Technical / Advisory	Annex 4: Qualification of IR spectrophotometers	Dec 2007
EDQM/OMCL TA	Technical / Advisory	Aide – Mémoire for environmental conditions and treatment of biological models	Jan 2011
EWDTs TA	Technical / Advisory	European Laboratory Guidelines for Legally Defensible Workplace Drug Testing  <i>Contact Person: R. Jonsson</i>	2002
EWDTs TA	Technical / Advisory	Drug and Alcohol Testing in Hair, Collection and Analysis	Aug 2010
EWDTs TA	Technical / Advisory	Guidelines for oral fluid	March 2011

### 2.6.2 Inspection Bodies - ISO/IEC 17020

Reference	Classification	Title	Date of publication	Date end transition (blank if not applicable)
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### 2.6.3 Certification Bodies ISO/IEC 17065 - Products

Reference	Classification	Title	Date of publication	Date end transition (blank if not applicable)
EA-6/02 M (rev.02)	Mandatory	EA Guidelines on the Use of EN 45011 and ISO/IEC 17021 for Certification to EN ISO 3834	June 2013	
EA-6/04 M (rev.00)	Mandatory	EA Guidelines on the Accreditation of Certification of Primary Sector Products by Means of Sampling of Sites	July 2011	

**2.6.4 Certification Bodies - ISO/IEC 17021-1 - Management Systems**

Reference	Classification	Title	Date of publication	Date end transition (blank if not applicable)
EA-7/04 M (rev.03)	Mandatory	Legal Compliance as a part of accredited ISO 14001: 2004 certification	May 2017	

**2.6.5 Verification Bodies - ISO 14065 - GHG**

Reference	Classification	Title	Date of publication	Date end transition (blank if not applicable)
EA-6/03 M (rev.04)	Mandatory	EA Document For Recognition of Verifiers under EU ETS Directive	Nov 2013	

**3 INTERNATIONAL DOCUMENTS APPLICABLE TO EA, EA MEMBERS AND EA MLA SIGNATORIES****3.1 ILAC DOCUMENTS** (all ILAC documents are available under <http://ilac.org/publications-and-resources/>)**3.1.1 ILAC POLICY DOCUMENTS (P-SERIES)**

Reference	Title	Applicable to EA as Region	Applicable to all	Applicable to the EA members signatories to ILAC/IAF	Date of publication	Date end transition (blank if not applicable)
ILAC-P4	ILAC Mutual Recognition Arrangement (Arrangement): Policy Statement			X	Feb 2016	
ILAC-P5	ILAC Mutual Recognition Arrangement (Arrangement)			X	Feb 2016	
ILAC-P8	ILAC Mutual Recognition Arrangement (Arrangement): Supplementary Requirements and Guidelines for the Use of Accreditation Symbols and for Claims of Accreditation Status by Accredited Laboratories and Inspection Bodies			X	Dec 2012	
ILAC-P9	ILAC Policy for Participation in Proficiency Testing Activities		X		June 2014	
ILAC-P10	ILAC Policy on the Traceability of Measurement Results		X		Jan 2013	
ILAC-P12	Harmonisation of ILAC Work with the Regions	X			April 2009	
ILAC-P13	Application of ISO/IEC 17011 for the Accreditation of Proficiency Testing Providers		X		Oct 2010	
ILAC-P14	ILAC Policy for Uncertainty in Calibration		X		Jan 2013	
ILAC-P15	Application of ISO/IEC 17020:2012 for the Accreditation of Inspection Bodies		X		Jul 2016	

**3.1.2 ILAC RULES DOCUMENTS (R-SERIES)**

Reference	Title	Applicable to EA as Region	Applicable to all	Applicable to the EA members signatories to ILAC/IAF	Date of publication	Date end transition (blank if not applicable)
ILAC-R4	Use of the ILAC logo and tagline			X	Oct 2016	
ILAC-R7	Rules for the Use of the ILAC MRA Mark			X	May 2015	

**3.1.3 ILAC GUIDANCE DOCUMENTS (G-SERIES)**

Reference	Title	Applicable to EA as Region	Applicable to all	Applicable to the EA members signatories to ILAC/IAF	Date of publication	Date end transition (blank if not applicable)
ILAC-G3	Guidelines for Training Courses for Assessors Used by Accreditation Bodies		X		Aug 2012	
ILAC-G18	Guidelines for the Formulation of Scopes of Accreditation for Laboratories			X	Apr 2010	
ILAC-G21	Cross Frontier Accreditation - Principles for Avoiding Duplication			X	Sept 2012	

**3.2 IAF/ILAC JOINT DOCUMENTS**

Reference	Title	Applicable to EA as Region	Applicable to all	Applicable to the EA members signatories to ILAC/IAF	Date of publication	Date end transition (blank if not applicable)
IAF/ILAC A1	IAF/ILAC MRAs: Evaluation of a Regional Group	X			Feb 2017	March 2019
IAF/ILAC A2	IAF/ILAC MRAs: Evaluation of a Single Accreditation Body	X			Feb 2017	March 2019
IAF/ILAC A3	IAF/ILAC Multi-Lateral Mutual Recognition Arrangements (Arrangements): Narrative	X			Jan 2013	

Framework for Reporting on the Performance  
of an Accreditation Body - A Tool for the  
Evaluation Process

IAF/ILAC A5	IAF/ILAC Multi-Lateral Mutual Recognition Arrangement (Arrangements): Application of ISO/IEC 17011:2004	X	Nov 2013
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### 3.3 IAF DOCUMENTS (all IAF documents are available under <http://www.iaf.nu//articles/Publications/6>)

#### 3.3.1 IAF POLICY DOCUMENTS (PL-SERIES)

Reference	Title	Applicable to EA as Region	Applicable to all	Applicable to the EA members signatories to ILAC/IAF	Date of publication	Date end transition (blank if not applicable)
IAF PL 1	Code of Conduct for Members of the IAF			X	Dec 2009	
IAF PL 6	Memorandum of Understanding			X	Feb 2016	
IAF-PL 8	Rules for the Use of the IAF Logo			X	April 2016	

#### 3.3.2 IAF MULTILATERAL RECOGNITION ARRANGEMENT DOCUMENTS (ML-SERIES)

Reference	Title	Applicable to EA as Region	Applicable to all	Applicable to the EA members signatories to ILAC/IAF	Date of publication	Date end transition (blank if not applicable)
IAF ML 1	Procedure for Exchange of Documentation among IAF MLA Accreditation Bodies			X	Jan 2016	
IAF ML 2	General Principles for Use of the IAF MLA Mark			X	May 2016	
IAF ML 3	Procedure on Responding to Inquiries on Multilateral Recognition Arrangement (MLA) Signatory Equivalence			X	Oct 2012	



IAF ML 4	Policies and Procedures for a MLA on the level of Single Accreditation Bodies and on the Level of Regional Accreditation Groups	X	May 2016
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### 3.3.3 IAF MANDATORY DOCUMENTS (MD-SERIES)

Reference	Title	Applicable to EA as Region	Applicable to all	Applicable to the EA members signatories to ILAC/IAF	Date of publication	Date end transition (blank if not applicable)
IAF MD 1	Certification of Multiple Sites Based on Sampling		X		Nov 2007	
IAF MD 2	Transfer of Accredited Certification of Management Systems		X		Dec 2007	
IAF MD 3	Advanced Surveillance and Recertification Procedures (ASRP)		X		Feb 2008	
IAF MD 4	Use of Computer Assisted Auditing Techniques ("CAAT") for Accredited Certification of Management Systems		X		May 2008	
IAF MD 5	Duration of QMS and EMS Audits		X		June 2015	
IAF MD 6	Application of ISO 14065:2013			X	March 2014	
IAF MD 7	Harmonization of Sanctions			X	Sept 2010	
IAF MD 8	Application of ISO/IEC 17011 in Medical Devices QMS (ISO 13485)		X		Jan 2015	
IAF MD 9	Application of ISO/IEC 17021 in Medical Devices QMS (ISO 13485)		X		Jan 2015	
IAF MD 10	Assessment of Certification Body Management of Competence in accordance with ISO/IEC 17021: 2011		X		Feb 2013	
IAF MD 11	IAF Mandatory Document for the Application of ISO/IEC 17021 for Audits of Integrated Management Systems		X		Dec 2013	

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IAF MD 12	Assessment of Certification Activities for Cross-Frontier Accreditation	X	Jan 2016	
IAF MD 13	Knowledge Requirements for Accreditation Body Personnel for Information Security Management Systems (ISO/IEC 27001)	X	Jan 2015	
IAF MD 14	Application of ISO/IEC 17011 in Greenhouse Gas Validation and Verification (ISO 14065:2013)	X	June 2014	
IAF MD 15	IAF Mandatory Document for the Collection of Data to Provide Indicators of Management System Certification Bodies' Performance	X	July 2014	
IAF MD 16	Application of ISO/IEC 17011 for the Accreditation of Food Safety Management Systems (FSMS) Certification Bodies	X	Jan 2015	
IAF MD 17	Witnessing Activities for the Accreditation of Management Systems Certification Bodies	X	Jan 2015	
IAF MD 18	Application of ISO/IEC 17021:2011 in the Service Management Sector (ISO/IEC 20000-1)	X	Jan 2015	
IAF MD 19	IAF Mandatory Document For The Audit and Certification of a Management System operated by a Multi-Site Organization (where application of site sampling is not appropriate)	X	March 2016	March 2018
IAF MD 20	Generic Competence for AB Assessors: Application to ISO/IEC 17011		X	May 2016 May 2018

## 4 INTERNATIONAL STANDARDS APPLICABLE TO EA, EA MEMBERS AND EA MLA SIGNATORIES

### 4.1 LEVEL 1 STANDARDS

Reference	Title	Date of publication	Date end transition (blank if not applicable)
EN ISO/IEC 17011	Conformity assessment -- General requirements for accreditation bodies accrediting conformity assessment bodies	2004	

### 4.2 LEVEL 3 STANDARDS

Reference	Title	Date of publication	Date end transition (blank if not applicable)
EN ISO/IEC 17025	General requirements for the competence of testing and calibration laboratories	2005	
EN ISO 15189	Medical laboratories -- Requirements for quality and competence	2012	1 March 2016
EN ISO/IEC 17020	Conformity assessment -- Requirements for the operation of various types of bodies performing inspection	2012	
EN ISO/IEC 17065	Conformity assessment -- Requirements for bodies certifying products, processes and services	2012	
EN ISO/IEC 17021-1	Conformity assessment -- Requirements for bodies providing audit and certification of management systems	2015	15 June 2017
EN ISO/IEC 17024	Conformity assessment – General requirements for bodies operating certification of persons	2012	
EN ISO 14065	Greenhouse gases -- Requirements for greenhouse gas validation and verification bodies for use in accreditation or other forms of recognition	2013	
EN ISO/IEC 17043	Conformity assessment -- General requirements for proficiency testing	2010	
ISO 17034	General requirements for the competence of reference material producers	2016	Nov 2019

**4.3 LEVEL 4 STANDARDS**

Reference	Title	Date of publication	Date end transition (blank if not applicable)
ISO/TS 22003	Food safety management systems -- Requirements for bodies providing audit and certification of food safety management systems	2013	15 Dec 2016
ISO/IEC 27006	Information technology -- Security techniques -- Requirements for bodies providing audit and certification of information security management systems	2015	30 Sept 2017
ISO 22870	Point-of-care testing (POCT) -- Requirements for quality and competence	2016	
ISO 15195	Laboratory medicine -- Requirements for reference measurement laboratories	2003	
CEN/TS 15675	Air quality. Measurement of stationary source emissions. Application of EN ISO/IEC 17025:2005 to periodic measurements	2007	
ISO/IEC TS 17021-2	Conformity assessment - Requirements for bodies providing audit and certification of management systems -- Part 2: Competence requirements for auditing and certification of environmental management systems	2016	Dec 2018
ISO/IEC TS 17021-3	Conformity assessment - Requirements for bodies providing audit and certification of management systems -- Part 3: Competence requirements for auditing and certification of quality management systems	2017	To be announced
ISO/IEC TS 17021-4	Conformity assessment – Requirements for bodies providing audit and certification of management systems -- Part 4: Competence requirements for auditing and certification of event sustainability management systems	2013	
ISO/IEC TS 17021-5	Conformity assessment -- Requirements for bodies providing audit and certification of management systems -- Part 5: Competence requirements for auditing and certification of asset management systems	2014	
ISO/IEC TS 17021-6	Conformity assessment -- Requirements for bodies providing audit and certification of management systems -- Part 6: Competence requirements for auditing and certification of business continuity management systems	2014	
ISO/IEC TS 17021-7	Conformity assessment -- Requirements for bodies providing audit and certification of management systems -- Part 7: Competence requirements for auditing and certification of road traffic safety management systems	2014	

ISO 50003 Energy management systems -- Requirements for bodies providing audit and certification of energy management systems 2014

#### 4.4 LEVEL 5 STANDARDS

Reference	Title	Date of publication	Date end transition (blank if not applicable)
ISO 14001	Environmental Management Systems – Requirements with guidance for use	2015	15 Sept 2018
ISO 9001	Quality management systems -- Requirements	2015	15 Sept 2018