MOLDAC REPUBLICA MOLDOVA	**NATIONAL ACCREDITATION CENTRE**

GENERAL CRITERIA FOR ACCREDITATION

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CONTENT

Note: The present document represents the English version of the Romanian document. In case of conflict the Romanian version will prevail.

1. PURPOSE

The purpose of this document is to describe the general criteria for accreditation of CABs by the National Accreditation Body MOLDAC, including criteria for the MOLDACs operation. At the same time, this document includes also MOLDACs policy regarding establishment of suitability of schemes and conformity assessment standards for the accreditation purposes.

2. SCOPE

Document is applied by MOLDAC personnel involved in accreditation process of CABs, as well as by all stakeholders.

3. REFERENCE DOCUMENTS

Reference documents are those listed at point 6 Description of activities of this document, including:

SR EN ISO / IEC 17000:2017 - Conformity assessment. Vocabulary and general principles.

RM Law no. 235 of 01.12.2011 on accreditation and conformity assessment activities, with subsequent amendments.

4. DEFINITIONS AND ABBREVIATIONS

4.1. Definitions

General criteria for accreditation of CABs - requirements stated in relevant normative documents such as international standards and guidelines for the operation of CABs.

4.2. Abbreviations

ONA	National Accreditation Body
CAB/OEC	Conformity Assessment Body
SM	Management System
LÎ	Testing Laboratory
LE	Calibration Laboratory
LVM	Metrological Verification Laboratory
LM	Medical Laboratory
OI	Inspection Body
OI/NDT	Inspection Body/ Nondestructive Testing
OCpr	Product Certification Body
OCsmc	Certification Body for Quality Management Systems
OCsmsa	Certification Body for food safety management systems
OCP	Personnel certification body
OPT	Organizer of proficiency testing and inter-laboratory comparisons
PT	Proficiency Testing
ILC	Inter-laboratory comparisons

5. ACCREDITATION SCHEMES. MOLDAC POLICY REGARDING ESTABLISHMENT OF ACCREDITAION SCHEMES

For each established accrditation scheme, MOLDAC evaluates the suitability of conformity assessment schemes at national level, as well as of the standards for the accreditation purpose, using I-03 Instruction, which was elaborated based on EA-1/22 document, and was endorsed by Technical Committee members.
For European conformity assessment schemes, MOLDAC makes sure that those were recognized at EA level, according to EA-1/22. For international conformity assessment schemes, MOLDAC makes sure that those were officaly approved by EA, ILAC or IAF.
MOLDAC commits to respect EA, ILAC or IAF decisions regarding the scheme, from the moment it becomes officially approved or withdrawn within this organizations.

MOLDAC established, and has competence, for following accreditation schemes:

5.1. Testing (level 2)

Conformity assessment standard: **SM EN ISO/IEC 17025:2017**[1] *(level 3)*

CAB scheme code	*CAB*	*Specific sectorial schemes and standards (level 4): normative documents used by accredited CABs (level 5)*
LI	Testing laboratory	Level 4: SM SR CEN/TS 15675:2012 Level 5: National legislation (Laws, GD, TR)

5.2. Testing: medical examinations (level 2)

Conformity assessment standard: **SM EN ISO 15189:2014** *(level 3)*

CAB scheme code	*CAB*	*Specific sectorial schemes and standards (level 4): normative documents used by accredited CABs (level 5)*
LM	Medical laboratory	Level 4: SM EN ISO 22870:2017 Level 5: National legislation (Laws, GD, TR)

5.3. Calibration (level 2)

Conformity assessment standard: **SM EN ISO/IEC 17025:2017**[2] *(level 3)*

CAB scheme code	*CAB*	*Specific sectorial schemes and standards (level 4): normative documents used by accredited CABs (level 5)*
LE	Calibration laboratory	Level 4: SM EN ISO 15195:2019 Level 5: National legislation (Laws, GD, TR)

5.4. Inspection (level 2)

Conformity assessment standard: **SM EN ISO/IEC 17020:2013** *(level 3)*

CAB scheme code	*CAB*	*Specific sectorial schemes and standards (level 4):*
OI	Inspection body	Level 5: National legislation (Laws, GD, TR)
LVM	Laboratory for metrological verification	Level 5: National legislation (Laws, GD, TR) RGML
OI NDT	Inspection body for non-destructive testing	Level 5: SM SR EN ISO 9712:2014, National legislation (Laws, GD, TR)

[1] During transition period will be considered SM SR EN SISO/IEC 17025:2006
[2] During transition period will be considered SM SR EN SISO/IEC 17025:2006

5.5. Product certification (level 2)

Conformity assessment standard: **SM EN ISO/IEC 17065:2013** *(level 3)*

CAB scheme code	*CAB*	*Specific sectorial schemes and standards (level 4):*	*Codes and standards or other normative documents used by accredited CABs, or private certification scheme (level 5):*
OCpr	Product certification body	National legislation harmonized with CE Regulations	National legislation (Laws, GD, TR)
OCprec	Organic products certification body	National legislation harmonized with CE Regulations (for example wirh Regulation (CE) 834/2007 for certification of organic products)	Law no.115/2003

5.6. Management systems certification (level 2)

Conformity assessment standard: **SM EN ISO/IEC 17021-1:2015** *(level 3)*

CAB scheme code	*CAB*	*Specific sectorial schemes and standards (level 4):*	*Codes and standards or other normative documents used by accredited CABs, or private certification scheme (level 5):*
OCsmc	Management systems certification bodies	SM ISO/IEC 17021-3:2017	SM SR EN ISO 9001:2015
OCsmsa		SM ISO/TS 22003:2014	SM SR EN ISO 22000:2006/ SM EN ISO 22000:2018

5.7 Certificare persoanelor (nivel 2)

Conformity assessment standard: **SM SR EN ISO/CEI 17024:2014** *(level 3)*

CAB scheme code	*CAB*	*Specific sectorial schemes and standards (level 3):*	*Codes and standards or other normative documents used by accredited CABs, or private certification scheme (level 5):*
OCP	Personnel certification body	SM SR EN ISO/CEI 17024:2014	National legislation (Laws, GD, TR)

6. DESCRIPTION OF ACTIVITIES

6.1. General criteria for National Accreditation Body MOLDAC from Republic of Moldova are defined in:

6.1.1 SM SR EN ISO / IEC 17011:2006 – “General requirements for accreditation bodies accrediting conformity assessment bodies”.

6.1.2 ISO/CEI 17011:2017 „Requirements for Accreditation Bodies accrediting Conformity Assessmnet Bodies”.

6.1.3 Regulation (EC) No 765/2008 of the European Parliament and of the Council of 9 July 2008 setting out the requirements for accreditation and market

surveillance relating to the marketing of products and repealing Regulation (EEC) No 339/93.

6.1.4 RM Law no. 235 of 01.12.2011 on accreditation and conformity assessment activities, with subsequent modification.

6.1.5 EA, ILAC, IAF documents according to EA/INF-01 List in force

- EA – 1/06 **A+AB:**2017	mandatory	EA Multilateral Agreement Criteria for signing Policy and procedure for development
- EA-1/13 A: 2016	Mandatory	EA's Relationship with Accreditation Bodies of Countries Not Being Members of the EU or EFTA
- EA – 1/17 S1 **A+AB** :2014	mandatory	Supliment 1 to EA-1/17, Criteria for Membership
- EA – 1/17 S3 **A** :2018	mandatory	EA supplement 3 to EA-1/17 Rules of Procedures. EA Procedure For the investigation and Resolution of Complaints and Appeals
- EA – 1/17-S5 **A+AB**:2018	mandatory	EA supplement 5 to EA-1/17, EA Rules of Procedure Levying of Membership Fees
- EA – 1/19 **A+AB**: 2016	mandatory	Rules for Use of the EA Logo
- EA-1/20-S1 **A+AB**:2016	mandatory	EA Supplement 1 to EA-1/20. Terms and Conditions for financial compensation from the operating/action grant to an EA Member Accreditation Body
- EA-1/21 **A+AB**:2008	mandatory	EA Internal Procedure for Liaison Activities
- EA – 1/22 **A**:2016	mandatory	EA Procedure and Criteria for the Evaluation of Conformity Assessment Schemes by EA Accreditation Body Members
- EA – 1/22 A-AB:2020	obligatoriu (cu aplicare din 14.04.2020)	EA Procedure and Criteria for the Evaluation of Conformity Assessment Schemes by EA Accreditation Body Members
- EA – 1/23 **A**:2019	mandatory	EA policy to speak with „One Voice"
- EA –2/02M:2019	mandatory	EA Procedure for the evaluation of a National Accreditation Body
- EA–2/13M :2019	mandatory	EA Cross Border Accreditation Policy and Procedure for Cross Border Cooperation

			between EA Members
-	EA–2/15M:2019	mandatory	EA Requirements for the Accreditation of Flexible Scopes
-	EA – 2/17M :2016	mandatory	EA Document on Accreditation for Notification purposes
-	**EA- 2**/17 **M**: 2020	obligatoriu (Cu aplicare din 14.04.2020)	EA Document on Accreditation for Notification Purposes
-	EA – 3/01M :2019	mandatory	EA conditions for the use of accreditation symbols, text reference to accreditation and MLA signatory status
-	ILAC G3:08/2012	guidelines	Guidelines for Training Courses for Assessors Used by Accreditation Bodies
-	ILAC G11:07/2006	guidelines	ILAC Guidelines on Qualifications & Competence of Assessors and Technical Experts
-	ILAC R4:10/2016	mandatory	Use of the ILAC logo and tagline
-	ILAC R7:05/2015	mandatory	Rules for the Use of the ILAC MRA Mark
-	IAF PL 8:2016	mandatory	Rules for the use of the IAF logo
-	IAF ML 2:2016	mandatory	General Principles on the Use of the IAF MLA Mark
-	IAF MD 20:2016	mandatory	Generic Competence for AB Assessors: Application to ISO/IEC 17011
-	IAF/ILAC A3:01/2018	mandatory	IAF/ILAC Multi-Lateral Mutual Recognition Arrangements (Arrangements): Template report for the peer evaluation of an Accreditation Body based on ISO/IEC 17011:2017
-	IAF/ILAC A5: 2013	mandatory	IAF/ILAC Multi-Lateral Mutual Recognition Arrangements (Arrangements): Application of ISO/IEC 17011:2004
-	ILAC-P8:03/ 2019	mandatory (implementation until march,2020)	ILAC Mutual Recognition Arrangement (Arrangement): Supplementary Requirements for the Use of Accreditation Symbols and for Claims of Accreditation Status by Accredited Conformity Assessment Bodies

6.2. General criteria for accreditation of the Testing Laboratories are defined in:

6.2.1 SM SR EN ISO/IEC 17025:2006 „General requirements for the competence of testing and calibration laboratories".

6.2.2 SM EN ISO/IEC 17025:2018 „General requirements for the competencce of Teting and Calibration laboratories".

6.2.3 EA, ILAC documents for Laboratory Accreditation Department, LÎ Section:

-	EA - 4/21:2018	informativ	Guidelines for the assessment of the appropriateness of small interlaboratory comparisons within the process of laboratory accreditation
-	ILAC P9:06/2014	mandatory	ILAC Policy for Participation in National and International Proficiency Testing Activities
-	ILAC-P10:01/2013	mandatory	ILAC Policy on Traceability of Measurement Results
-	ILAC G8:09/2019	guidelines	Guidelines on Decision Rules and Statements of Conformity
-	ILAC G18:04/2010	guidelines	Guidelines for the Formulation of Scopes of Accreditation for Laboratories
-	ILAC G21:09/2012	guidelines	Cross Frontier Accreditation – Principles for Cooperation

6.2.4 EA, ILAC documents for Testing Laboratories:

-	EA – 4/09:2017	guidelines	Accreditation for Sensory Testing Laboratories (previously EAL-G16)
-	EA – 4/14:2003	informative	Selection and Use of Reference Materials
-	EA – 4/16:2003	guidelines	EA Guidelines on the Expression of Uncertainty in Quantitative testing
-	EA – 4/18:2010	informative	Guidance on the level and frequency of proficiency testing participation
-	EA - 4/21:2018	informativ	Guidelines for the assessment of the appropriateness of small interlaboratory comparisons within the process of laboratory accreditation
-	EA-4/23:2016	informativ	The Assessment and Accreditation of Opinions and Interpretations using ISO/IEC 17025:2017
-	ILAC-P10:01/2013	mandatory	ILAC Policy on Traceability of Measurement

			Results
-	ILAC G8:09/2019	guidelines	Guidelines on Decision Rules and Statements of Conformity
-	ILAC G19:08/2014	guidelines	Modules in a Forensic Science Process
-	ILAC G24:2007	guidelines	Guidelines for the determination of calibration intervals of measuring instruments
-	ILAC-P8:03/ 2019	mandatory (with aplication from march, 2020)	ILAC Mutual Recognition Arrangement (Arrangement): Supplementary Requirements for the Use of Accreditation Symbols and for Claims of Accreditation Status by Accredited Conformity Assessment Bodies

6.2.5 MOLDAC documents placed on the website www.acreditare.md.

-	Policy P-02	Policy on use of proficiency testings and of other interlaboratory coparisons in accreditation process
-	Policy P-03	Policy on measurement of traceability
-	Policy P-04	Policy on handling of nonconformities
-	Policy P-05	Policy cross-border accreditation
-	Policy P-07	Policy on treating the objections of CABs regarding assessment team members and observers
-	Policy P-08	Policy and ryles on the use of accreditation symbols, of ILAC-MRA combined mark and references to accreditation
-	RA	Rules for Accreditation
-	DR-LÎ/LE-01	Guidelines for acreditation of testing and calibration laboratories
-	DI-LÎ/LE-10	Informative document regarding transition to the new version of ISO/IEC 17025:2017

6.3. General criteria for accreditation of Calibration Laboratories are defined in:

6.3.1 SM SR EN ISO/CEI 17025:2006 „General requirements for the competence of testing and calibration laboratories”.

6.3.2 SM EN ISO/IEC 17025:2018 „General requirements for the competencce of Teting and Calibration laboratories”.

6.3.3 EA, ILAC documents for Laboratories Accreditation Department, LE section:

-	EA - 4/21:2018	informativ	Guidelines for the assessment of the appropriateness of small interlaboratory comparisons within the process of laboratory accreditation
-	ILAC P9:06/2014	mandatory	ILAC Policy for Participation in National and International Proficiency Testing Activities
-	ILAC-P10:01/2013	mandatory	ILAC Policy on Traceability of Measurement Results
-	ILAC G8:09/2019	guidelines	Guidelines on Decision Rules and Statements of Conformity
-	ILAC G18:04/2010	guidelines	Guidelines for the Formulation of Scopes of Accreditation for Laboratories

6.3.4 EA, ILAC documents for Calibration Laboratories:

-	EA – 4/02:2013	mandatory	Expressions of the Uncertainty of Measurements in Calibration (including supplement 1 to EA-4/02)
-	EA - 4/21:2018	informativ	Guidelines for the assessment of the appropriateness of small interlaboratory comparisons within the process of laboratory accreditation
-	ILAC-P10:01/2013	mandatory	ILAC Policy on Traceability of Measurement Results
-	ILAC-P14:01/2013	mandatory	ILAC Policy for uncertainty in Calibration
-	ILAC G8:09/2019	guidelines	Guidelines on Decision Rules and Statements of Conformity
-	ILAC G24:2007	guidelines	Guidelines for the determination of calibration intervals of measuring instruments
-	ILAC-P8:03/ 2019	mandatory (with application from march, 2020)	ILAC Mutual Recognition Arrangement (Arrangement): Supplementary Requirements for the Use of Accreditation Symbols and for Claims of Accreditation Status by Accredited Conformity Assessment Bodies

6.3.5 MOLDAC documents placed on the website www.acreditare.md.

-	Policy P-02	Policy on use of proficiency testings and of other interlaboratory coparisons in accreditation process

- Policy P-03		Policy on measurement of traceability
- Policy P-04		Policy on handling of nonconformities
- Policy P-05		Policy cross-border accreditation
- Policy P-07		Policy on treating the objections of CABs regarding assessment team members and observers
- Policy P-08		Policy and ryles on the use of accreditation symbols, of ILAC-MRA combined mark and references to accreditation
- RA		Rules for Accreditation
- DR-LÎ/LE-01		Guidelines for acreditation of testing and calibration laboratories
- DI-LÎ/LE-10		Informative document regarding transition to the new version of ISO/IEC 17025:2017

6.4. General criteria for accreditation of the Medical Laboratories are defined in:

6.4.1. SM SR EN ISO 15189:2014 „Medical labratories. Requirements for quality and competence".

6.4.2. EA, ILAC documents for Laboratory Accreditation Department, LM section:

-	EA - 4/17:2008	mandatory	EA Position Paper on the description of scopes of accreditation of medical laboratories
-	EA – 4/20:2015	guidelines	Guidance for the Assessment of Laboratories against EN ISO 15189 and 22870
-	EA - 4/21:2018	informativ	Guidelines for the assessment of the appropriateness of small interlaboratory comparisons within the process of laboratory accreditation
-	EA-4/23:2016	informativ	The Assessment and Accreditation of Opinions and Interpretations using ISO/IEC 17025:2017
-	ILAC P9:06/2014	mandatory	ILAC Policy for Participation in National and International Proficiency Testing Activities
-	ILAC-P10:01/2013	mandatory	ILAC Policy on Traceability of Measurement Results
-	ILAC G26:11/2018	guidelines	Guidance for the implementation of the Medical Laboratory Accreditation System

6.4.3. EA, ILAC documents for Medical Laboratories:

-	EA – 4/14:2003	informative	Selection and use of reference materials
-	EA – 4/16:2003	guidelines	EA guidelines on the expression of uncertainty in quantitative testing
-	EA-4/17:2008	mandatory	EA Position Paper on the description of scopes of accreditation of medical laboratories
-	EA – 4/18:2010	informative	Guidance on the level and frequency of proficiency testing participation
-	EA - 4/21:2018	informativ	Guidelines for the assessment of the appropriateness of small interlaboratory comparisons within the process of laboratory accreditation
-	ILAC-P10:01/2013	mandatory	ILAC Policy on Traceability of Measurement Results
-	ILAC-P8:03/ 2019	mandatory (with application from march, 2020)	ILAC Mutual Recognition Arrangement (Arrangement): Supplementary Requirements for the Use of Accreditation Symbols and for Claims of Accreditation Status by Accredited Conformity Assessment Bodies
-	ILAC-P8:03/ 2019	guidelines	Modules in a Forensic Science Process
-	ILAC G24:2007	guidelines	Guidelines for the determination of calibration intervals of measuring instruments

6.4.4. MOLDAC documents placed on the website www.acreditare.md.

-	Policy P-02	Policy on use of proficiency testings and of other interlaboratory coparisons in accreditation process
-	Policy P-03	Policy on measurement of traceability
-	Policy P-04	Policy on handling of nonconformities
-	Policy P-05	Policy cross-border accreditation
-	Policy P-07	Policy on treating the objections of CABs regarding assessment team members and observers
-	Policy P-08	Policy and ryles on the use of accreditation symbols, of ILAC-MRA combined mark and references to accreditation
-	RA	Rules for Accreditation
-	DR-LM-03	Guidelines for acreditation of medical laboratories

6.5. General criteria for accreditation of the Inspection Bodies and Metrological Verifications Laboratories are defined in:

6.5.1. SM SR EN ISO/CEI 17020:2013 “Conformity assessment. Requirements for operation of different types of bodies which performs the inspection”.

6.5.2. EA, ILAC documents for Laboratory Accreditation Direction/ Accreditation Directorate Certification and Inspection Bodies, Sections OI/NDT and LVM:

- EA - 4/21:2018	informativ	Guidelines for the assessment of the appropriateness of small interlaboratory comparisons within the process of laboratory accreditation
- ILAC P9:06/2014	mandatory	ILAC Policy for Participation in National and International Proficiency Testing Activities
- ILAC-P10:01/2013	mandatory	ILAC Policy on Traceability of Measurement Results
- ILAC-P15:07/2016	mandatory	Application of ISO/IEC 17020:2012 for the Accreditation of Inspection Bodies
ILAC P15:05/2020 -	obligatoriu (cu aplicare din noiembrie 2021)	Application of ISO/IEC 17020:2012 for the Accreditation of Inspection Bodies
- IAF MD 8:2017	mandatory (with application from 09.06.2018)	Application of ISO/IEC 17011:2004 4 in the Field of Medical Device Quality Management Systems (ISO 13485)
- ILAC-G:28:07/2018	mandatory	Guideline for the Formulation of Scopes of Accreditation for Inspection Bodies

6.5.3. EA, ILAC documents for Inspection Bodies and Metrological Verification Laboratories:

- EA - 4/21:2018	informativ	Guidelines for the assessment of the appropriateness of small interlaboratory comparisons within the process of laboratory accreditation
- ILAC-P10:01/2013	mandatory	ILAC Policy on Traceability of Measurement Results
- ILAC-P15:07/2016	mandatory	Application of ISO/IEC 17020:2012 for the Accreditation of Inspection Bodies
- ILAC P15:05/2020 -	obligatoriu (cu aplicare din noiembrie 2021)	Application of ISO/IEC 17020:2012 for the Accreditation of Inspection Bodies

-	ILAC G19:08/2014	guidelines	Modules in a Forensic Science Process
-	ILAC G24:2007	guidelines	Guidelines for the determination of calibration intervals of measuring instruments
-	ILAC-G:27:07/2019	guidelines	Guidance on measurements performed as part of an inspection process
-	ILAC-G:27:06/2017	guidelines	Guidance on measurements performed as part of an inspection process
-	ILAC-P8:03/ 2019	mandatory (with application from march, 2020)	ILAC Mutual Recognition Arrangement (Arrangement): Supplementary Requirements for the Use of Accreditation Symbols and for Claims of Accreditation Status by Accredited Conformity Assessment Bodies

6.5.4. MOLDAC documents placed on the website www.acreditare.md.

-	Policy P-02	Policy on use of proficiency testings and of other interlaboratory coparisons in accreditation process
-	Policy P-03	Policy on measurement of traceability
-	Policy P-04	Policy on handling of nonconformities
-	Policy P-05	Policy cross-border accreditation
-	Policy P-07	Policy on treating the objections of CABs regarding assessment team members and observers
-	Policy P-08	Policy and ryles on the use of accreditation symbols, of ILAC-MRA combined mark and references to accreditation
-	RA	Rules for Accreditation
-	DR-LVM-02	Guidelines for accreditation of metrological verification laboratories
-	DR-OI-07	Guidelines for accreditation of inspection bodies
-	DR-OI/NDT-08	Guidelines for accreditation of inspection bodies/ nondestructive testings

6.6. General criteria for accreditation of the Products Certification Bodies defined in:

6.6.1. SM SR EN ISO/CEI 17065:2013 „Conformity assessment. Requiremetns for bodies that certify products, processes and services”.

6.6.2. EA, IAF documents for Certification and Inspection Bodies Accreditation Department:

- EA - 3/12:2013	mandatory	EA Policy for Accreditation of Organic Production Certification
- IAF MD 7:2010	mandatory	Harmonization of Sanctions
- IAF MD 12:2016	mandatory	Accreditation Assessment of Conformity Assessment Bodies with Activities in Multiple Countries
- IAF ML 2:2016	mandatory	General Principles on the Use of the IAF MLA Mark

6.6.3. EA, IAF documents for Product Certification Bodies:

- EA 6/04:2011	– mandatory	EA Guidelines on the accreditation of certification of primary sector products by means of sampling of sites

6.6.4 MOLDAC documents placed on the website www.acreditare.md

- Policy P-02	Policy on use of proficiency testings and of other interlaboratory coparisons in accreditation process
- Policy P-03	Policy on measurement of traceability
- Policy P-04	Policy on handling of nonconformities
- Policy P-05	Policy cross-border accreditation
- Policy P-07	Policy on treating the objections of CABs regarding assessment team members and observers
- Policy P-08	Policy and ryles on the use of accreditation symbols, of ILAC-MRA combined mark and references to accreditation
- RA	Rules for Accreditation
- DR-OCpr-04	Guidelines for accreditaiton of product certification bodies
- I-03	Criteria for assessment of conformity assessment schemes
- CS-OCprec-01	Specific criteria for accreditation of ecologic product certification bodies
- CS-OCpr-02	Specific requirements for accreditation of protected origin and geographical indications vine products certification bodies

6.7. General criteria for accreditation of the Management Systems Certification Body System Management of Quality defined in:

6.7.1. SM SR ISO/CEI 17021-1:2015 “Conformity assessment. Requirements for bodies providing audit and certification of management systems. Part 1: Requirements”

6.7.2. SM SR ISO/CEI 17021-3:2017 ”Conformity assessment. Requirements for bodies providng audit and certification of management systems Part 3: Cerințe de Competence requirements for auditing and certification of quality management systems”.

6.7.3. SM SR EN ISO 9001:2015 ”Quality management system. Requirements”.

6.7.4. EA, IAF document for Certification and Inspection Bodies Accreditation Department:

- IAF MD 2:2017	mandatory	IAF mandatory document for transfer of management systems certification issued under accreditation
- IAF MD 7:2010	mandatory	Harmonization of Sanctions
- IAF MD 10:2013	mandatory	IAF Mandatory Document for Assessment of Certification Body Management of Competence in Accordance with ISO/IEC 17021:2011
- IAF MD 12:2016	mandatory	Accreditation Assessment of Conformity Assessment Bodies with Activities in Multiple Countries
- IAF MD 15:2014	mandatory	Mandatory Document for the Collection of Data to Provide Indicators of Management System Certification Bodies’ Performance
- IAF MD 17:2019	mandatory	Witnessing Activities for the Accreditation of Management Systems Certification Bodies
- IAF ID 1:2014	informative	IAF Informative Document for QMS Scopes of Accreditation
- IAF ML 2:2016	mandatory	General Principles on the Use of the IAF MLA Mark

6.7.5. EA, IAF documents for Quality management system Certification Bodies

- IAF MD 1:2018	mandatory	IAF Mandatory Document for the Audit and Certification of Management System Operated by a Multi-Site Organization
- IAF MD 2:2017	mandatory	IAF mandatory document for transfer of management systems certification issued under

			accreditation
-	IAF MD 3:2008	mandatory	Advanced Surveillance and Recertification Procedures (ASRP)
-	IAF MD 4:2018	mandatory	IAF Mandatory Document for the Use of Computer Assissted Auditing Techniques (CAAT) for Accredited Certification of Management Sistems
-	IAF MD 5:2019	mandatory	Determination of audit time of quality, enviromental, and occupational health &safety management systems
-	IAF MD 11:2019	obligatoriu (with aplication from 17.01.2020)	IAF Mandatory Document for the Application of ISO/IEC 17021-1 for Audits of Integrated Management Systems
-	IAF ID 1:2014	informative	IAF Informative Document for QMS Scopes of Accreditation

6.7.6. MOLDAC documents placed on the website www.acreditare.md:

-	Policy P-04	Policy on handling of nonconformities
-	Policy P-05	Policy cross-border accreditation
-	Policy P-07	Policy on treating the objections of CABs regarding assessment team members and observers
-	RA	Rules for Accreditation
-	DR-OCsmc-05	Guidelines for accreditation of management system certification bodies
-	DI-OCsmc-09	Informative document regarding transition to ISO/IEC 17021-3:2017 standard
-	DI-OCsmc-07	Informative document regarding transition to ISO 9001:2015 standard

6.8. General criteria for accreditation of the Certification Body Food Safety Management Systems defined in:

6.8.1 SM SR ISO/CEI 17021-1:2015 “Conformity assessment. Requirements for bodies providing audit and certification of management systems. Part 1: Requirements”

6.8.2 SM ISO/TS 22003:2014 “Food safety management systems. Requirements for bodies providing audit and certification of food safety management systems”

6.8.3 SM EN ISO 22000:2018 ”Food Safety management system. Requirements for any organization from food chain”.

6.8.4 EA, IAF documents for Certification and Inspection Bodies Accreditation Department:

- IAF MD 2:2017	mandatory	IAF mandatory document for transfer of management systems certification issued under accreditation
- IAF MD 7:2010	mandatory	Harmonization of Sanctions
- IAF MD 10:2013	mandatory	IAF Mandatory Document for Assessment of Certification Body Management of Competence in Accordance with ISO/IEC 17021:2011
- IAF MD 12:2016	mandatory	Accreditation Assessment of Conformity Assessment Bodies with Activities in Multiple Countries
- IAF MD 15:2014	mandatory	Mandatory Document for the Collection of Data to Provide Indicators of Management System Certification Bodies’ Performance
- IAF MD 16:2015	mandatory	Application of ISO/IEC 17011 for the Accreditation of Food Safety Management Systems (FSMS) Certification Bodies

6.8.5 EA, IAF documents for Food Safety management system Certification Bodies:

- IAF MD 2:2017	mandatory (with aplication from 15.06.2018)	IAF Mandatory Document for transfer of management sistems certifications issued under accreditation
- IAF MD 11:2019	mandatory (with aplication from 17.01.2020)	IAF Mandatory Document for the Application of ISO/IEC 17021-1 for Audits of Integrated Management Systems
- IAF ML 2:2016	obligatoriu	General Principles on the Use of the IAF MLA Mark

6.8.6 MOLDAC documents placed on the website www.acreditare.md:

- Policy P-04	Policy on handling of nonconformities
- Policy P-05	Policy cross-border accreditation
- Policy P-07	Policy on treating the objections of CABs regarding assessment team members and observers
- Policy P-08	Policy and ryles on the use of accreditation symbols, of ILAC-MRA combined mark and references to accreditation

-	RA	Rules for Accreditation
-	DR-OCsmsa-06	Guidelines for accreditation of food safety management system certification bodies

6.9 6.8. General criteria for accreditation of the Personnel certification body defined in::

6.9.1 SM SR EN ISO/IEC 17024:2014 „ Conformity assessment – General requirements for bodies operating certification of persons".

6.9.2 SM ISO/IEC TS 17027:2014 „Conformity assessment — Vocabulary related to competence of persons used for certification of persons"

6.9.3 EA, IAF documents for Certification and Inspection Bodies Accreditation Department:

-	IAF MD 7:2010	obligatoriu	Harmonozation of Sanctions
-	IAF MD 12:2016	obligatoriu	Accreditation Assessment of Conformity Assessment Bodies with Activities in Multiple Countries

6.9.3 EA, IAF documents for Personnel Certification Body:
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6.9.4 MOLDAC documents placed on the website www.acreditare.md:

-	Policy P-04	Policy on handling of nonconformities
-	Policy P-05	Policy cross-border accreditation
-	Policy P-07	Policy on treating the objections of CABs regarding assessment team members and observers
-	Policy P-08	Policy and ryles on the use of accreditation symbols, of ILAC-MRA combined mark and references to accreditation
-	RA	Rules for Accreditation
-	DR-OCP-09	Guidelines for accreditation Personnel certification body

7. SYNTHESIS OF CHANGES

There were included modification on the following pages 1, 3, 4, 5, 8, 11, 19.