ROSATOM

ROSATOM STATE CORPORATION FOR NUCLEAR ENERGY

APCS certification

Company: RASU JSC

Moscow

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Contents

European Union. General Requirements.

International Electrotechnical Commission – IEC

International Organization for Standardization, ISO

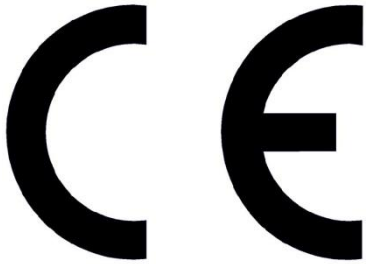
IEEE
(Institute of Electrical and Electronics Engineers)

National standardization bodies:
Hungary - MSZT; Finland - SFS, Germany – DIN,
France - AFNOR, etc.

Basic Standards

Nuclear Safety Standards
IEC 61226:2010
IEC 61513:2013
IEC 60987:2015
IEC 60880:2009
IEC 62138:2009
IEC 62340:2010
IEC/IEEE 60780-323:2016
IEC 61500:2011
IEC 61508:2010
IEC 62556:2014
IEC 61227:2008
IEC 61225:2005
IEC 60709:2004

General industrial directives of the European Parliament
2014/30/EU – Electromagnetic compatibility
2014/35/EU – Low voltage equipment
2014/32/EU – Instrumentation
2014/34/EU (ATEX) – Explosion-Proof Equipment



Hierarchy of requirements as exemplified by the PAKS II Project

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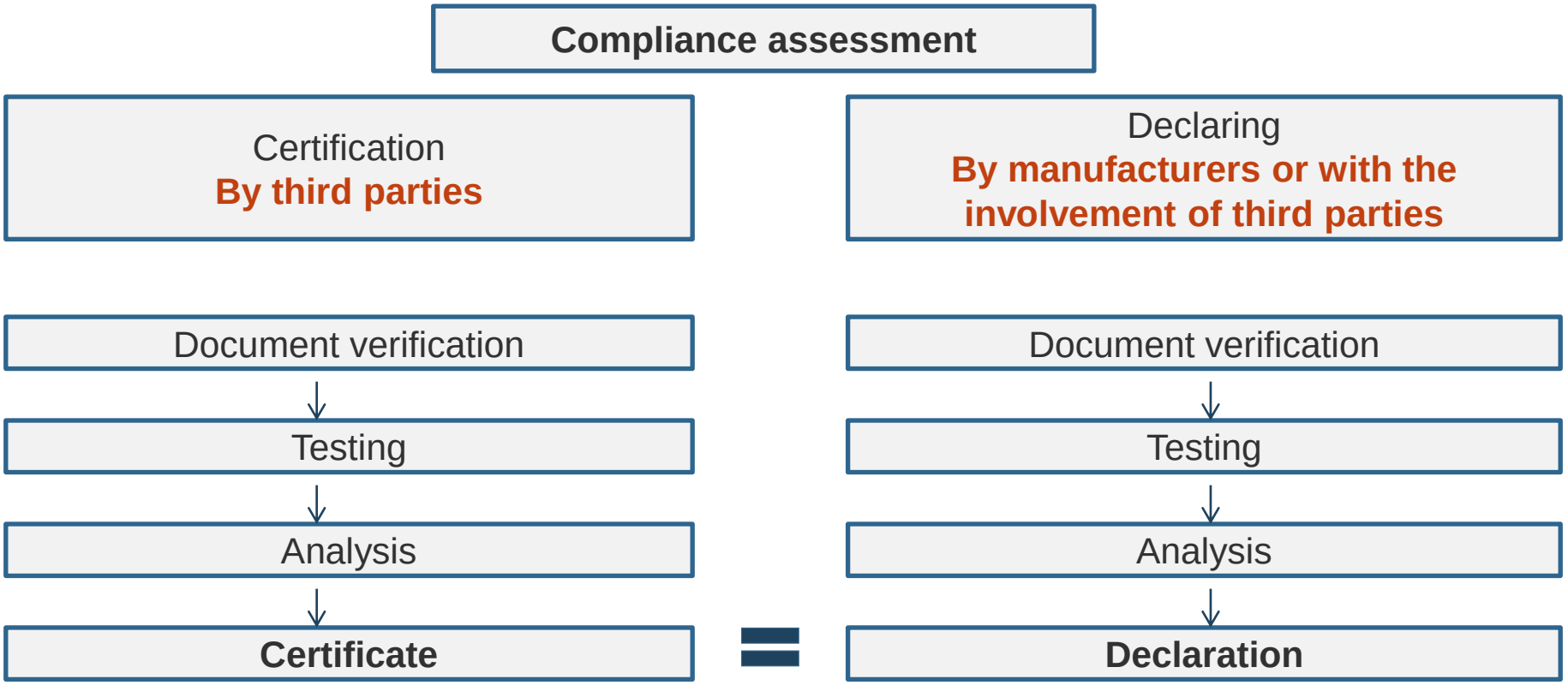
Compliance assessment. Certification / Declaring

Conformity assessment concept

Conformity assessment is the process of confirming that the requirements established for the product, process, service, system, person or body have been met.



Conformity confirmation procedures. Solution 768/2008/EC

<table>
<tr><th>Manufacturing</th><th>Designing</th></tr>
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In-house production control</td></tr>
<tr><td>Module C
The conformity of products to the sample tested</td><td rowspan="4">Module B
Testing product samples according to the rules of the European Union with third parties involved</td></tr>
<tr><td>Module D
Availability of a quality assurance system</td></tr>
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Availability of a quality assurance system and product inspection</td></tr>
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Availability of an engineering, production, testing, and control assurance system</td></tr>
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Conformity assessment stages

Stage 1 Assessing the conformity of the assessment scheme and the list of standards to the EU Directives.

Stage 2 Bringing the engineering documentation in line with the requirements of the European Union.

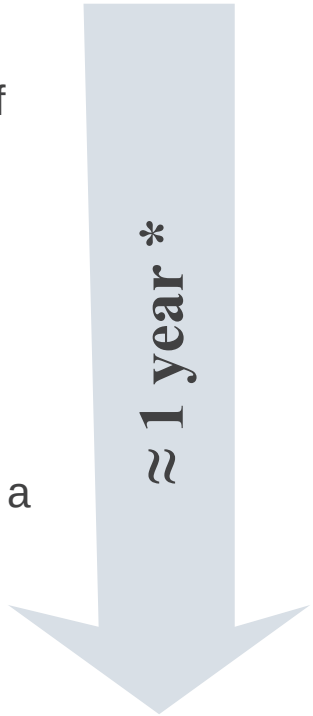
Stage 3 Conducting tests by the manufacturer and/or using the resources of an accredited laboratory (if applicable).

Stage 4 Checking the documentation by a notified certification body.

Stage 5 Conducting audit of production and the quality management system by a notified certification body (if applicable).

Stage 6 Assessing conformity of products in the form of certification and declaring.

Stage 7 Conducting supervisory audit (once per 1 to 3 years) - confirmation of conformity of products to the declared standards (if applicable).



* Provided there is a complete set of documentation for the supplied equipment by the beginning of certification and selection at Phase 3 of a laboratory already accredited in the EU

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Criteria for selecting a body for conformity assessment.

An independent conformity assessment body authorized to carry out conformity assessment shall be accredited in accordance with **ISO/IEC 17065** for this type of product

An independent inspection body authorized to carry out inspection check-up shall be accredited in accordance with **ISO/IEC 17020** for this type of product

An independent testing laboratory authorized to conduct product tests shall comply with the requirements of **ISO/IEC 17025** for this type of product

Certification bodies and testing laboratories accredited in the European Union

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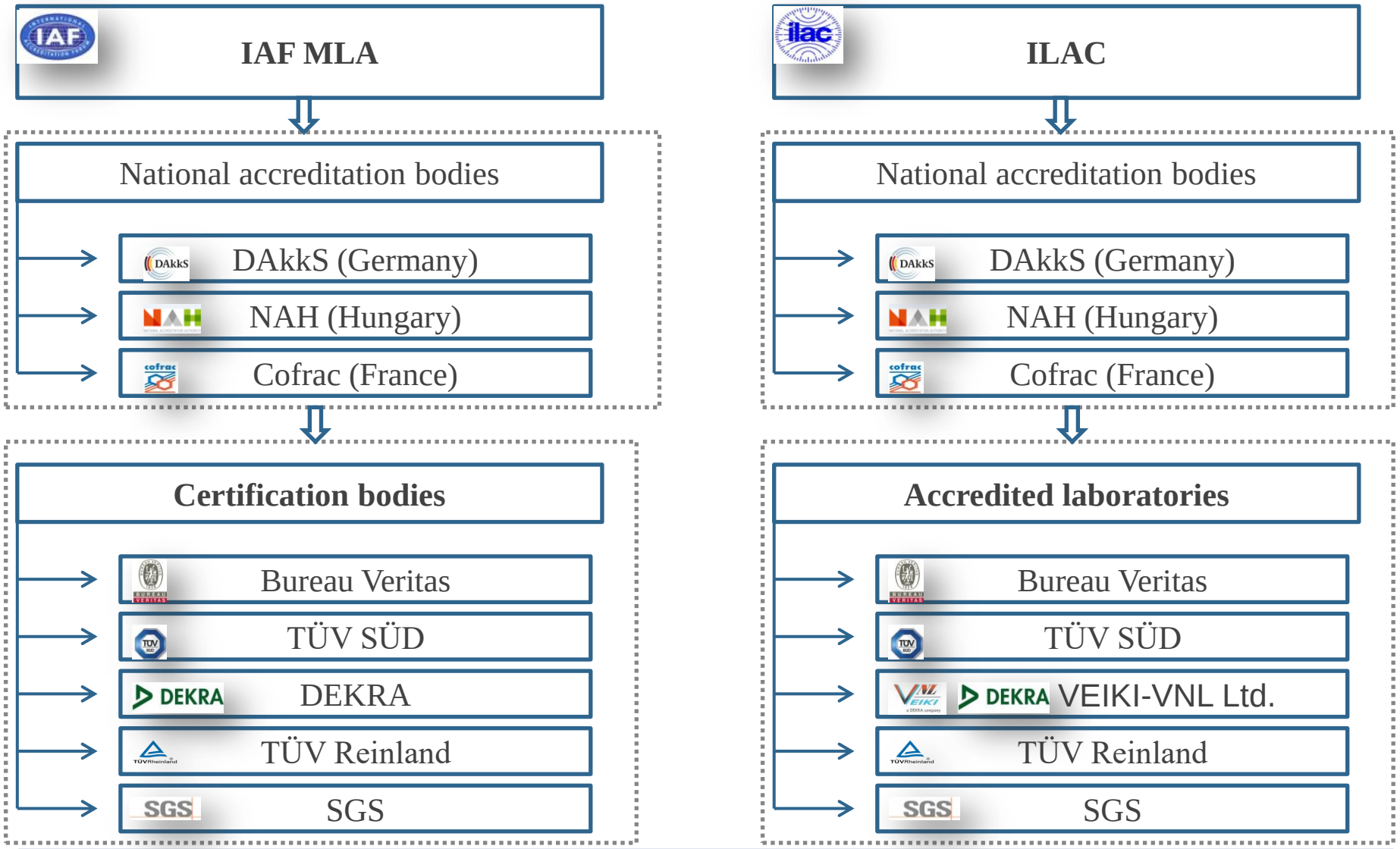

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Thank you for your time!