

**DESIGNATION OF A CONFORMITY ASSESSMENT BODY (CAB) FROM (*THIRD COUNTRY*) PURSUANT TO A MUTUAL RECOGNITION AGREEMENT
BETWEEN THE EUROPEAN UNION AND (*THIRD COUNTRY CONCERNED*)**

Date:

From: (*Third Country concerned*)

[*Name and address of Authority responsible for the designation of the body*]

To: European Commission
DG for Trade
Unit TRADE/G/3
B-1049 Brussels

1. References

Sectoral annex:

Directive/...../.....

2. Organisation name, acronym, address, telephone/fax, e-mail, web address

Name:

Abbreviated name:

Address:

Town/City:

Country:

Telephone:

Fax:

Company e-mail:

Web site:

Contact person(s)

3. Period of validity of the designation

- Unlimited
- Valid until (dd/mm/yyyy)

4. Technical qualifications of the body (accreditation or other official authorisation):

[*Body's name*] meets the requirements in the annex [*annex to the directive laying down the minimum criteria*].

What criteria have been used to assess the competence of the body?

- Legal personality
- Independence of organisation (3rd - party)
- Independence of personnel
- Professional integrity
- Technical knowledge (tasks/procedures)
- Technical training / knowledge of essential requirements
- Impartiality
- Liability insurance
- Professional secrecy

Competence assessment performed by:

- Designating Authority: [*name*]

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- Nationally recognised accreditation body: [*name*]

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Accreditation/assessment in accordance with:

- EN 45001 - EN ISO/IEC 17025 - testing laboratories
- EN 45004 - EN ISO/IEC 17020 – inspection
- EN 45011 - product certification
- EN 45012 - EN ISO/IEC 17021 - certification of quality systems
- EN 45013 - EN ISO/IEC 17024 - certification of personnel
- EN ISO/IEC 17025 - testing and calibration laboratories
- Other (please specify)

Note: The standards indicated must meet the requirements of the relevant conformity assessment procedures.

Please attach and list here the supporting documentation such as relevant accreditation scope if applicable (*pdf format preferable*)

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Information on surveillance procedures (including frequency)

- Arrangements for monitoring the body:
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- Review due on: (dd/mm/yyyy)

5. Designation procedure used to determine the competence of the proposed CAB to apply conformity assessment requirements and procedures identified in the legislative, regulatory and administrative provisions of the EU
(For example documentary review and/or details of on-site peer assessments).

6. Tasks performed by the body

Product/product range	Procedures/Modules	Annexes/Articles of the legislation

Is the NB accredited/assessed to perform all these procedures?

- Yes/No

Are any of these procedures sub-contracted to other bodies?

- Yes/No

Is the body a member of a Notified Bodies' co-ordination group?

- Yes/No
- Name of NB coordination group