ICS 03.120.20

Atbilstības novērtēšana. Piegādātāja atbilstības deklarācija. 1.daļa: Vispārīgās prasības (ISO/IEC 17050-1:2004, koriģētā versija 2007-06-15)

LVS EN ISO/IEC 17050-1

2010. g. 16.septembris

Aizstāj LVS EN ISO/IEC 17050-1:2005 A un LVS EN ISO/IEC 17050-1:2005 L

Conformity assessment - Supplier's declaration of conformity - Part 1: General requirements (ISO/IEC 17050-1:2004, corrected version 2007-06-15)

EIROPAS STANDARTS EN ISO/IEC 17050-1:2010

PĀRŅEMTS LATVIJAS STANDARTA STATUSĀ

Nacionālais priekšvārds

Latvijas standarts LVS EN ISO/IEC 17050-1:2010 "Atbilstības novērtēšana. Piegādātāja atbilstības deklarācija. 1.daļa: Vispārīgās prasības (ISO/IEC 17050-1:2004, koriģētā versija 2007-06-15)" ir identisks Eiropas standartam EN ISO/IEC 17050-1:2010 "Conformity assessment - Supplier's declaration of conformity - Part 1: General requirements (ISO/IEC 17050-1:2004, corrected version 2007-06-15)".

Eiropas standarts bez pārveidojumiem tā saturā pārņemts nacionālā standarta statusā.

Lappuses: EN-2; ISO/IEC-6
Cenas grupa: E

Standartizācijas birojs (LVS) ir Latvijas nacionālā standartizācijas institūcija un viens no tās uzdevumiem ir nodrošināt juridiskās un fiziskās personas ar Latvijas standartiem. LVS pārstāv Latvijas viedokli Eiropas un starptautiskajās standartizācijas organizācijās. LVS ir tiesības izdot un publicēt Latvijas standartus. Latvijas standartus aizsargā Latvijas Republikas Autortiesību likums, starptautisko vienošanos normas un Eiropas un starptautisko standartizācijas organizāciju noteikumi autortiesību jomā.

Standartu piemērošana ir brīvprātīga. Ministru kabinets var noteikt obligāti piemērojamus standartus. Standartu lietošana novērš nesaskaņas, jo tie satur viennozīmīgas prasības. Standarti balstās uz jaunākajiem zinātnes un tehnikas sasniegumiem un veicina tautsaimniecības attīstību. Standartu lietošana nodrošina cilvēka dzīvības, veselības un vides aizsardzību, produktu un sistēmu saderību, veicina preču un pakalpojumu brīvu kustību.

Standarti nav mācību līdzekļi. Standartu lietotājam ir jābūt profesionālām zināšanām, lai pareizi lietotu standartus un atbildētu par savas darbības sekām.

Standartus var iegādāties **www.lvs.lv** vai LVS Standartu fondā:

K. Valdemāra ielā 157, Rīgā, LV-1013

Tālrunis/fakss 67362250

IF@lvs.lv

EUROPEAN STANDARD

NORME EUROPÉENNE

EUROPÄISCHE NORM

EN ISO/IEC 17050-1

April 2010

ICS 03.120.20

Supersedes EN ISO/IEC 17050-1:2004

English version

Conformity assessment - Supplier's declaration of conformity - Part 1: General requirements (ISO/IEC 17050-1:2004, corrected version 2007-06-15)

Évaluation de la conformité - Déclaration de conformité du fournisseur - Partie 1: Exigences générales (ISO/IEC 17050-1:2004, version corrigée 2007-06-15)

Konformitätsbewertung - Konformitätserklärung von Anbietern - Teil 1: Allgemeine Anforderungen (ISO/IEC 17050-1:2004, korrigierte Fassung 2007-06-15)

This European Standard was approved by CEN on 13 March 2010.

CEN and CENELEC members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard the status of a national standard without any alteration. Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the CEN Management Centre or to any CEN and CENELEC member.

This European Standard exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN and CENELEC member into its own language and notified to the CEN Management Centre has the same status as the official versions.

CEN and CENELEC members are the national standards bodies and national electrotechnical committees of Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.

CEN Management Centre:
Avenue Marnix 17, B-1000 Brussels

CENELEC Central Secretariat:
Avenue Marnix 17, B-1000 Brussels

Ref. No. EN ISO/IEC 17050-1:2010 E

Foreword

The text of ISO/IEC 17050-1:2004, corrected version 2007-06-15 has been prepared by Committee on conformity assessment (CASCO) of the International Organization for Standardization (ISO) and the International Electrotechnical Commission (IEC) and has been taken over as EN ISO/IEC 17050-1:2010 by Technical Committee CEN/CLC/TC 1 “Criteria for conformity assessment bodies” the secretariat of which is held by BSI.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by October 2010, and conflicting national standards shall be withdrawn at the latest by October 2010.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. CEN [and/or CENELEC] shall not be held responsible for identifying any or all such patent rights.

This document supersedes EN ISO/IEC 17050-1:2004.

According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland and the United Kingdom.

Endorsement notice

The text of ISO/IEC 17050-1:2004, corrected version 2007-06-15 has been approved by CEN as a EN ISO/IEC 17050-1:2010 without any modification.

Conformity assessment — Supplier's declaration of conformity —

Part 1:
General requirements

1 Scope

This part of ISO/IEC 17050 specifies general requirements for a supplier's declaration of conformity in cases where it is desirable, or necessary, that conformity of an object to the specified requirements be attested, irrespective of the sector involved. For the purposes of this part of ISO/IEC 17050, the object of a declaration of conformity can be a product, process, management system, person or body.

This part of ISO/IEC 17050 does not define any particular object for the declaration of conformity.

Instead of "supplier's declaration of conformity", the term "declaration of conformity" can be used when appropriate.

2 Normative references

The following referenced documents are indispensable for the application of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

ISO/IEC 17000:2004, *Conformity assessment — Vocabulary and general principles*

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO/IEC 17000 apply.

NOTE 1 "Supplier's declaration of conformity" is a "declaration" as defined in ISO/IEC 17000, i.e. first-party attestation.

NOTE 2 To avoid any confusion with attestation by certification bodies, the term "self-certification" is deprecated and should not be used.

4 Purpose of the declaration of conformity

The purpose of the declaration is to give assurance of conformity of the identified object to specified requirements to which the declaration refers, and to make clear who is responsible for that conformity and declaration. A supplier's declaration of conformity may be used alone or in conjunction with another conformity assessment procedure for regulatory or non-regulatory purposes.

5 General requirements

The issuer (issuing organization or person) of a declaration of conformity shall be responsible for issuing, maintaining, extending, reducing, suspending or withdrawing the declaration and the conformity of the object to the specified requirements.

The declaration of conformity shall be based on results of an appropriate type of conformity assessment activity (e.g. testing, measurement, auditing, inspection or examination) carried out by one or more first, second or third parties. Conformity assessment bodies involved, where applicable, should consult relevant International Standards, Guides and other normative documents.

Where a declaration of conformity is for a group of products of a similar type, it shall cover each individual product of the group. Where a declaration of conformity is for similar products delivered over a period of time, it shall cover each product as delivered or accepted.

It is recommended, as good conformity assessment practice, that the person reviewing the conformity assessment results be different from the signatory.

6 Contents of the declaration of conformity

6.1 The issuer of the declaration of conformity shall ensure that the declaration contains sufficient information to enable the recipient of the declaration of conformity to identify the issuer of the declaration, the object of the declaration, the standards or other specified requirements with which conformity is declared, and the person signing for and on behalf of the issuer of the declaration of conformity.

As a minimum, the declaration of conformity shall contain the following:

a) unique identification of the declaration of conformity;

b) the name and contact address of the issuer of the declaration of conformity;

c) the identification of the object of the declaration of conformity (e.g. name, type, date of production or model number of a product, description of a process, management system, person or body, and/or other relevant supplementary information);

d) the statement of conformity;

e) a complete and clear list of standards or other specified requirements, as well as the selected options, if any;

f) the date and place of issue of the declaration of conformity;

g) the signature (or equivalent sign of validation), name and function of the authorized person(s) acting on behalf of the issuer;

h) any limitation on the validity of the declaration of conformity.

6.2 Additional supporting information may be provided to relate the declaration to the conformity assessment results on which it is based, for example:

a) the name and address of any conformity assessment body involved (e.g. testing or calibration laboratory, inspection body, certification body);

b) reference to relevant conformity assessment reports, and the date of the reports;

c) reference to any management systems involved;

d) reference to the accreditation documents of conformity assessment bodies involved where the scope of accreditation is relevant to the declaration of conformity;

e) reference to the existence of associated supporting documentation, such as that described in ISO/IEC 17050-2;

f) additional information regarding certificates, registrations or marks that have been obtained;

g) other activities or programmes of the conformity assessment body (e.g. membership in an agreement group).

References in the documentation to conformity assessment results shall not misrepresent their applicability nor mislead the recipient of the declaration of conformity.

7 Form of declaration of conformity

See Annex A for an example of a declaration of conformity. The declaration of conformity may be in hardcopy, electronic media, or any other suitable medium.

8 Accessibility

A copy of the declaration of conformity may be included in other documentation, such as a statement, catalogue, invoice, user's instructions or website, relevant to the object of the declaration of conformity.

9 Product marking

If any marking is placed on the product to indicate the existence of a declaration of conformity, such marking shall be in such a format that it will not be confused with any certification mark. Such marking shall be traceable to the declaration of conformity.

10 Continuing validity of the declaration of conformity

10.1 The issuer of the declaration of conformity shall have procedures in place to ensure the continued conformity of the object, as delivered or accepted, with the stated requirements of the declaration of conformity.

10.2 The issuer of the declaration of conformity shall have procedures in place to re-evaluate the validity of the declaration of conformity, in the event of

a) changes significantly affecting the object’s design or specification,

b) changes in the standards to which conformity of the object is stated,

c) changes in the ownership or structure of management of the supplier, if relevant, or

d) relevant information indicating that the object may no longer conform to the specified requirements.

Annex A
(informative)

Supplier's declaration of conformity

A.1 Guidance to complete the form of declaration of conformity

NOTE Numbers 1) to 7) refer to the form shown in A.2.

1) Every declaration of conformity should be uniquely identified.

2) The responsible issuer should be unequivocally specified. For large organizations, it may be necessary to specify operational groups or departments.

3) a) The “object” should be unequivocally described so that the declaration of conformity may be related to the object in question.

3) b) For mass-produced products, it is not necessary to give individual serial numbers. In such cases it is sufficient to give the name, type, model number, etc.

4) For products, an alternative conformity statement may be: “As delivered, the object of the declaration described above is in conformity with the requirements of the following documents”.

5) Requirements documents should be listed with their identification numbers, titles and dates of issue.

6) Text should appear here only if any limitation on the validity of the declaration of conformity and/or any additional information are given. The latter information may, for example, correspond to 6.2 or may make reference to related product marking in accordance with Clause 9. Such product marking or other indication (e.g. on the product) may be an attachment to the declaration of conformity.

7) Full name and function of the signing person(s) authorised by the issuer's management to sign on its behalf should be given. The number of signatures, or equivalent, included will be the minimum determined by the legal form of the issuer’s organization.

A.2 Example of form of declaration of conformity

Supplier's declaration of conformity (in accordance with ISO/IEC 17050-1)

1) **No.** .....

2) **Issuer's name:** .....

Issuer's address: .....

.....

3) **Object of the declaration:** .....

.....

.....

4) **The object of the declaration described above is in conformity with the requirements of the following documents:**

	Documents No.	Title	Edition/Date of issue
5)	.....	.....	.....
	.....	.....	.....
	.....	.....	.....

Additional information:

6) .....

.....

.....

Signed for and on behalf of:

.....

.....

(Place and date of issue)

7) ....

(Name, function) (Signature or equivalent authorized by the issuer)

Bibliography

[1] ISO 9000:2000, *Quality management systems — Fundamentals and vocabulary*

[2] ISO 19011:2002, *Guidelines for quality and/or environmental management systems auditing*

[3] ISO/IEC 17020:1998, *General criteria for the operation of various types of bodies performing inspection*

[4] ISO/IEC 17021:2006, *Conformity assessment — Requirements for bodies providing audit and certification of management systems*

[5] ISO/IEC 17024:2003, *Conformity assessment — General requirements for bodies operating certification of persons*

[6] ISO/IEC 17025:2005, *General requirements for the competence of testing and calibration laboratories*

[7] ISO/IEC 17040:2005, *Conformity assessment — General requirements for peer assessment of conformity assessment bodies and accreditation bodies*

[8] ISO/IEC 17050-2:2004, *Conformity assessment — Supplier's declaration of conformity — Part 2: Supporting documentation*

[9] ISO/IEC Guide 65:1996, *General requirements for bodies operating product certification systems*