BS EN 50581:2012



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http: Technical documentation for the assessment of electrical and electronic products with respect to the restriction of hazardous substances

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National foreword

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des produits électriques et électroniques par rapport à la restriction des substances dangereuses

Beurteilung von Elektro- und Elektronikgeräten hinsichtlich der Beschränkung gefährlicher Stoffe

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Foreword

This document (EN 50581:2012) has been prepa	ared by C	LC/TC 111X "Envi	ronment".
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 This document (EN 50581:2012) has been prepared. The following dates are fixed: latest date by which this document has to be implemented at national level by publication of an identical national standard or by endorsement latest date by which the national standards conflicting with this document have to be withdrawn 	(dop)	2013-07-16 10 2013-07-16 2015-07-16	ges

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

This document has been prepared under a mandate given to CENELEC by the European Commission and the European Free Trade Association, and supports essential requirements of EU Directive 2011/65/EU.

For the relationship with EU Directive see informative Annex ZZ, which is an integral part of this document.

Introduction

Certain substances contained in electrical and electronic products are restricted by legislation and/or customer specifications. Manufacturers of final products therefore need to be able demonstrate that their products meet the applicable substance restrictions.

For those restrictions that apply at the "homogeneous material" level, it is productical for manufacturers of complex products to undertake their own testing of all materials contained in the final assembled product. Instead, manufacturers work with the suppliers to manage compliance and compile technical documentation as evidence of compliance. This approach is well recognised by both industry and enforcement authorivity.

The aim of this European Standard is to specify the technical documentation that the manufacturer needs to compile in order to recharge compliance with the applicable substance restrictions. In this way, this European Standard supports Directive 2011/65/EU of the European Parliament and of the Souncil of 8 June 2011 on the restriction of the use of certain hazardous substances in platition and electronic equipment (RoHS).

This European Standard can also find an application in demonstrating conformity to other substance regulations worldwide.

Scope 1

This European Standard specifies the technical documentation that the manufacturer needs to compile in order to declare compliance with the applicable substance restrictions.

The documentation of the manufacturer's management system is outside the scope of this European Standard. **2 Normative references** The following documents, in whole or in part, are normatively deterenced in this document and are indispensable for its application. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies. NN

EN 62321:2009, Electrotechnical products — Determination of levels of six regulated substances (lead, mercury commum, hexavalent chromium, polybrominated biphenyls, Determination of levels of six regulated polybrominated diphenyl ethers) (IEC 62321:2008)

NOTE EN 62321 will be replaced by a series of standards designated EN 62321-x.

3 Terms and definitions

For the purposes of this document, the following terms and definitions apply.

3.1

restricted substance

substance which is limited in its use in a product, part or material

3.2

manufacturer

natural or legal person who manufactures a product or has a product designed or manufactured, and markets that product under his name or trademark

[SOURCE: Regulation (EC) N° 765/2008 or Decision N° 768/2008/EC]

Note 1 to entry: In certain circumstances, an importer or distributor is considered a manufacturer for the purposes of Directive 2011/65/EU.

3.3

supplier

organisation that provides the manufacturer with materials, parts and/or sub-assemblies

Technical documentation 4

4.1 Overview

The manufacturer shall compile technical documentation to demonstrate that electrical and electronic products comply with substance restrictions (see 4.2 and 4.3).

4.2 Content of the technical documentation

The technical documentation shall include at least the following elements:

a general description of the product;

NOTE 1 Directive 2011/65/EU specifies 11 product categories. The product category is one of the factors that determines which exemptions apply.

- documents for materials, parts, and/or sub-assemblies (see 4.3);
- information showing the relationship between the technical documents identified in 4.3 and the corresponding materials, parts and/or sub-assemblies in the product;

- list of harmonized standards and/or other technical specifications that have been used to establish the technical documents identified in 4.3, or to which such documents refer.

NOTE 2 Annex A describes the relationship between Article 7(b) of Directive 2011/65/EU, Module And Decision 768/2008/EC, this European Standard, and the technical documentation.
4.3 Information on materials, parts, and/or sub-assemblies
4.3.1 Tasks to be undertaken by the manufacturer
The manufacturer shall undertake the following four tasks:
determine the information needed (see 4, 3.0)
collect the information (see 4.3.3):

- evaluate the information with bard to its quality and trustworthiness and decide whether to include it in the technical documentation (see 4.3.4);
- ensure that the techn a documentation remains valid (see 4.3.5).

Figure 1 (informative) shows the process to create the technical documentation:



Figure 1 — Schematic representation of process to create the technical documentation

4.3.2 Determine the information needed

The types of technical documents (see 4.3.3) that are required for materials, parts and/or subassemblies shall be based on the manufacturer's assessment of:

- a) the probability of restricted substances being present, in materials, parts or subassemblies, and
- b) the trustworthiness of the supplier.

Materials that are added during the production process (such as solder, paint, adhesives) shall also be considered as part of the assessment.

When undertaking the assessment described in point a), the manufacturer may apply technical judgement, as some substances are unlikely to be contained in certain materials (e.g. organic substances in metals). Such technical judgement could be based on technical information available via the electrical/electronic industry, or a literature investigation of the materials/parts used in electrical/electronic products.

NOTE 1 Additional information that can be used when undertaking the assessment described in points (a) and (b) includes:
material types typically used in the part or sub-assembly;
historical likelihood of restricted substances being present in exprematerial type;
historical experience with the supplier organization;
results of previous supplier inspections or audit.

NOTE 2 The assessment and its associated procedures can form part of a quality management system or equivalent.

4.3.3 Collecting informat

As a result of the manufacturer's assessment, the following documents on materials, parts, and/or sub-assemblies shall be collected:

- a) Supplier declarations and/or contractual agreements, such as:
 - Supplier declarations, confirming that the restricted substance content of the material, part, or sub-assembly is within the permitted levels and identifying any exemptions that have been applied;
 - Signed contracts confirming that the manufacturer's specification for the maximum content of restricted substances in a material, part, or sub-assembly is fulfilled.

Such declarations or agreements shall cover a specific material, part and/or subassembly, or a specific range of materials, parts and/or sub-assemblies.

and/or

- b) Material Declarations:
 - Material declarations providing information on specific substance content and identifying any exemptions that have been applied.

NOTE 1 The use of standards for such declarations helps ensure consistent and costeffective flow of information throughout the supply chain. EN 62474 "Material declaration for products of and for the electrotechnical industry" describes the procedure, content, and form relating to material declaration. Other specifications for material declarations are also used in industry today.

and/or

c) Analytical test results:

- Analytical test results using the methods described or referenced in EN 62321.

NOTE 2 EN 62321 will be replaced by a series of standards designated EN 62321-x.

4.3.4 Evaluation of information

The manufacturer shall establish procedures that shall be used to evaluate the documents described in 4.3.3 in order to determine their quality and trustworthiness.

NOTE 1 IEC/TR 62476 provides a framework for the use of internationally accepted standards, tools and practices to evaluate electrical and electronic products with respect to restricted substances.

The manufacturer shall evaluate, in accordance with these procedures, the source and content of each document received in order to determine whether or not the material, part, or sub-assembly meets the specified substance restrictions.

NOTE 2 The substance restrictions specified in Directive 2011/65/EU apply at the homogeneous material level.

This evaluation will enable the manufacturer to decide whether the documents provide sufficient evidence of compliance to justify their inclusion in the technical documentation. If a particular document is:

- considered to be of sufficient quality and trustworthiness, then it shall be included the technical documentation;
- not considered to be of sufficient quality or trustworthiness, then ufacturer actions are necessary – particulations include requesting additional information from the supplier or uncertaining his own substance analysis.
 Review of the technical documentation nanufacturer shall:
 perform a periodic review of the documents contained in the technical documentation to ansure that they are full relid.

4.3.5 Review of the technical documentation

The manufacturer shall:

- to ensure that they are still lid:
- ensure that the technical documentation reflects any changes to materials, parts or sub-assemblies in accordance with 4.3.3.

NOTE Directive 2011/65/EU requires that "manufacturers ensure that procedures are in place for series production to remain in conformity. Changes in product design or characteristics and changes in the harmonised standards or in technical specifications by reference to which conformity of EEE is declared shall be adequately taken into account".



(informative)



Figure A.1 — Relationship between Directive 2011/65/EU and the technical documentation covered by this standard

Annex ZZ

(informative)

Coverage of Essential Requirements of EU Directives

s.com This European Standard has been prepared under a mandate given to C by the European Commission and the European Free Trade Association Article its scope the standard covers all relevant essential requirements as gives of Directive 7 2011/65/EU.

Compliance with this standard provides one machs conformity with the specified essential requirements of the Directive concerned. WARNING: Other requirements and other Directives may be applicable to the products falling within the scope of this standard.

the scope of this standard.

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IEC/TR 62476:2010, Guidance for evaluation of products with restrictions in electrical and electronic products

products — exercision of restricted substance use IEC/PAS 62596:2009, Electrotechnical products -Sampling procedure — Guidelines

EN ISO 9001, Quality systems

agement systems — Requirements with guidance for use EN ISO 14001, Environn (ISO 14001)

Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment

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

Decision No 768/2008/EC of the European Parliament and of the Council of 9 July 2008 on a common framework for the marketing of products, and repealing Council Decision 93/465/EEC

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