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Annex ZA (informative) Relationship of this European Standard with the Construction Products Directive ZA.1 Clauses of this European Standard addressing the provisions of EU Construction Products Directive This European Standard has been prepared under Mandates M101 “External and internal doors and windows, roof

Annex ZA (informative) Relationship of this European ...

Annex ZZ (informative) Relationship between this European Standard and the safety requirements of Directive 2001/95/EC aimed to be covered. This European Standard has been prepared under a Commission’s standardization request [Full reference to the request “M/xxx”] to provide one voluntary means of conforming to the safety requirements of Commission Decision (EU/EC) No [XXX/YYYY] of ...

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Once this standard is cited in the Official Journal of the European Union under that [Directive] / [Regulation] / [Decision] / […],

compliance with the normative clauses of this standard given in Table […] confers, within the limits of the scope of this standard, a presumption of conformity with the corresponding [essential] / [interoperability] / […] requirements of that [Directive ...

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Once this standard is cited in the Official Journal of the European Union (OJEU), under Directive 2014/28/EU, compliance with the normative clauses of this standard given in Table ZA.1 confers, within the limits of the scope of this standard, a presumption of conformity with the corresponding essential requirements of that Directive 2014/28/EU, and associated EFTA regulations.

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Annex ZA (informative) Relationship of this European Standard with Regulation (EU) No.305/2011 (When applying this standard as a harmonized standard under Regulation (EU) No. 305/2011, manufacturers and Member States are obliged by this regulation to use this Annex) ZA.1 Scope and relevant characteristics

Implementation of the Construction Products Regulation ...

!For relationship with EU Directive(s), see informative Annex ZA, which is an integral part of this document." This standard is part of the EN 115 series of standards: “Safety of escalators and moving walks”. !deleted text" According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following

Safety of escalators and moving walks

For relationship with EU Directive(s), see informative Annex ZA, which is an integral part of this document. The main changes compared t o the previous edition are: a) the appropriate method for determination of bacterial filtration efficiency (BFE) provided in vitro in Annex B has been updated;

BS EN 14683:2019

Annex ZA (informative) Relationship between this European Standard and the essential requirements of EU Directive 2014/68/EU (PEI)) aimed to be covered This European Standard has been prepared under a Commission's standardization request M/071 "Mandate to CEN for standardization in the field of pressure equipment" to provide one voluntary

KM C754e-20161215102853

For relationship with EU Directive(s), see informative Annex ZA, which is an integral part of this standard. According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Czech

EUROPEAN STANDARD EN 81-1

Annex Za Informative Relationship Of annex. ZA.2.2 Procedure according to system 3 For products falling under system 3, the

tasks for the approved body and the manufacturer related to the initial type test and the factory production control are described in Table ZA.3.

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For relationship with EU Directive 97/23/EC, see informative Annex ZA, which is an integral part of this document. The main changes are: new definitions: manufacturer , intermediary , product specification ; reduction of the number of inspection documents: type 2.3 of the previous edition has been deleted;

Metallic products - Types of inspection documents

Annex ZZ (informative) Relationship between this European standard and the safety objectives of Directive 2014/35/EU [2014 OJ L96] aimed to be covered. ... Annex ZA proposed by EC_2015-05-29_LVD specifities included_marked-up ...

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For relationship with EU Directive(s), see informative Annexes ZA, B and C, which are an integral part of this document. The following referenced documents are indispensable for the application of this document.

Packaging for terminally sterilized medical devices ...

For relationship with EU Directive(s), see informative Annex ZA, B, C or D, which is an integral part of this document. FprEN 16584-1:2015 (E) 5 . Introduction . This document is part of a suite of four ‘Design for PRM Use’ standards that have in total nine parts:

Railway Applications Design for PRM Use - General ...

Annex ZA(informative)Relationship of this European Standard and the essential requirements of Regulation (EU) 2019/1009

making available on the market of EU fertilising products aimed to be covered . This European Standard has been prepared under a standardization request M/564 annexed to Commission