



SLOVENSKI STANDARD

SIST EN 61331-2:2014

01-december-2014

Nadomešča:
SIST EN 61331-2:2002

**Sredstva za zaščito pred rentgenskim sevanjem pri medicinski diagnostiki - 2. del:
Prosojne zaščitne plošče (IEC 61331-2:2014)**

Protective devices against diagnostic medical X-radiation - Part 2: Translucent protective plates

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Dispositifs de protection radiologique contre les rayonnements X pour diagnostic médical
- Partie 2: Plaques translucides de protection radiologique

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Ta slovenski standard je istoveten z: EN 61331-2:2014

ICS:

11.040.50	Radiografska oprema	Radiographic equipment
13.280	Varstvo pred sevanjem	Radiation protection

SIST EN 61331-2:2014

en

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EUROPEAN STANDARD

EN 61331-2

NORME EUROPÉENNE

EUROPÄISCHE NORM

October 2014

ICS 11.040.50

Supersedes EN 61331-2:2002

English Version

Protective devices against diagnostic medical X-radiation - Part 2: Translucent protective plates (IEC 61331-2:2014)

Dispositifs de protection radiologique contre les
rayonnements X pour diagnostic médical - Partie 2: Plaques
translucides de protection radiologique
(CEI 61331-2:2014)

Strahlenschutz in der medizinischen Röntgendiagnostik -
Teil 2: Durchsichtige Schutzplatten
(IEC 61331-2:2014)

This European Standard was approved by CENELEC on 2014-06-11. 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-CENELEC Management Centre or to any 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 CENELEC member into its own language and notified to the CEN-CENELEC Management Centre has the same status as the official versions.

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European Committee for Electrotechnical Standardization
Comité Européen de Normalisation Electrotechnique
Europäisches Komitee für Elektrotechnische Normung

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

Foreword

The text of document 62B/937/FDIS, future edition 2 of IEC 61331-2, prepared by SC 62B, "Diagnostic imaging equipment", of IEC TC 62, "Electrical equipment in medical practice " was submitted to the IEC-CENELEC parallel vote and approved by CENELEC as EN 61331-2:2014.

The following dates are fixed:

- latest date by which the document has to be implemented at national level by publication of an identical national standard or by endorsement (dop) 2015-04-24
- latest date by which the national standards conflicting with the document have to be withdrawn (dow) 2017-06-11

This document supersedes EN 61331-2:2002.

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Endorsement notice

The text of the International Standard IEC 61331-2:2014 was approved by CENELEC as a European Standard without any modification.

IEC 60601-2-17:2013

NOTE

Harmonised as EN 60601-2-17:2014.

[SIST EN 61331-2:2014](https://standards.iteh.ai/catalog/standards/sist/7bbe0da0-7c04-427e-8e46-9a4e41398ca5/sist-en-61331-2-2014)

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Annex ZA (normative)

Normative references to international publications with their corresponding European publications

The following documents, in whole or in part, are normatively referenced 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.

NOTE 1 When an International Publication has been modified by common modifications, indicated by (mod), the relevant EN/HD applies.

NOTE 2 Up-to-date information on the latest versions of the European Standards listed in this annex is available here: www.cenelec.eu.

<u>Publication</u>	<u>Year</u>	<u>Title</u>	<u>EN/HD</u>	<u>Year</u>
IEC 60601-1	2005	Medical electrical equipment -- Part 1: General requirements for basic safety and essential performance	EN 60601-1	2006
			+EN 60601-1:2006/corrigendum Mar. 2010	2010
			+AC	2014
			+A11	2011
+A1	2012		+A1	2013
IEC 60601-1-3	2008	Medical electrical equipment -- Part 1-3: General requirements for basic safety and essential performance - Collateral Standard: Radiation protection in diagnostic X-ray equipment	EN 60601-1-3	2008
			+EN 60601-1-3:2008/corrigendum Mar. 2010	2010
			+A1	2013
			+AC	2014
+A1	2013		FprEN 60601-2-8	2010
IEC 60601-2-8	2010	Medical electrical equipment -- Part 2-8: Particular requirements for basic safety and essential performance of therapeutic X-ray equipment operating in the range 10 kV to 1 MV		
IEC 61331-1	2014	Protective devices against diagnostic medical X-radiation -- Part 1: Determination of attenuation properties of materials	EN 61331-1	2014
ISO 3534-1	2006	Statistics_ - Vocabulary and symbols_ - Part_1: General statistical terms and terms used in probability	-	-
IEC/TR 60788	2004	Medical electrical equipment - Glossary of defined terms	-	-
ISO/IEC Guide 99	2007	International vocabulary of metrology - Basic and general concepts and associated terms (VIM)	-	-

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IEC 61331-2

Edition 2.0 2014-05

INTERNATIONAL STANDARD

NORME INTERNATIONALE

Protective devices against diagnostic medical X-radiation –
Part 2: Translucent protective plates
(standards.iteh.ai)

Dispositifs de protection radiologique contre les rayonnements X pour
diagnostic médical –
Partie 2: Plaques translucides de protection radiologique

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INTERNATIONAL ELECTROTECHNICAL COMMISSION

**PROTECTIVE DEVICES AGAINST
DIAGNOSTIC MEDICAL X-RADIATION –**
Part 2: Translucent protective plates**FOREWORD**

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International Standard IEC 61331-2 has been prepared by subcommittee 62B: Diagnostic imaging equipment, of IEC technical committee 62: Electrical equipment in medical practice.

This second edition cancels and replaces the first edition of IEC 61331-2, published in 1994. It constitutes a technical revision. This second edition has been adapted to apply to the present technology.

The essential changes and extensions are:

extension of scope to cover all kinds of TRANSLUCENT PROTECTIVE PLATES and all kinds of RADIATION QUALITIES and GAMMA RADIATION;

removal of definition and requirements for TRANSLUCENT PROTECTIVE PLATES for visual imaging;
changes of requirements concerning geometrical accuracy and optical quality;

changes of requirements concerning determination of LEAD EQUIVALENT and minimal thickness;