



SLOVENSKI STANDARD

SIST EN 61223-3-4:2002

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Evaluation and routine testing in medical imaging departments - Part 3-4: Acceptance tests - Imaging performance of dental X-ray equipment

Evaluation and routine testing in medical imaging departments -- Part 3-4: Acceptance tests - Imaging performance of dental X-ray equipment

Bewertung und routinemäßige Prüfung in Abteilungen für medizinische Bildgebung -- Teil 3-4: Abnahmeprüfungen - Leistungsmerkmale zur Bildgebung von zahnärztlichen Röntgeneinrichtungen

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Essais d'évaluation et de routine dans les services d'imagerie médicale -- Partie 3-4: Essais d'acceptation - Performance d'imagerie des appareils de radiographie dentaire

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Evaluation and routine testing in medical imaging departments
Part 3-4: Acceptance tests
Imaging performance of dental X-ray equipment
(IEC 61223-3-4:2000)

Essais d'évaluation et de routine dans les services d'imagerie médicale
Partie 3-4: Essais d'acceptation
Performance d'imagerie des appareils de radiographie dentaire
(CEI 61223-3-4:2000)

Bewertung und routinemäßige Prüfung in Abteilungen für medizinische Bildgebung
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Leistungsmerkmale zur Bildgebung von zahnärztlichen Röntgeneinrichtungen
(IEC 61223-3-4:2000)

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Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the Central Secretariat 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 Central Secretariat has the same status as the official versions.

CENELEC members are the national electrotechnical committees of Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Netherlands, Norway, Portugal, Spain, Sweden, Switzerland and United Kingdom.

CENELEC

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

Central Secretariat: rue de Stassart 35, B - 1050 Brussels

Foreword

The text of document 62B/393/FDIS, future edition 1 of IEC 61223-3-4, 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 was approved by CENELEC as EN 61223-3-4 on 2000-06-01.

The following dates were fixed:

- latest date by which the EN has to be implemented at national level by publication of an identical national standard or by endorsement (dop) 2001-03-01
- latest date by which the national standards conflicting with the EN have to be withdrawn (dow) 2003-06-01

Annexes designated "normative" are part of the body of the standard.
Annexes designated "informative" are given for information only.
In this standard, annexes A and ZA are normative and annexes B and C are informative.
Annex ZA has been added by CENELEC.

Endorsement notice

The text of the International Standard IEC 61223-3-4:2000 was approved by CENELEC as a European Standard without any modification.

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In the official version, for Bibliography, the following note has to be added for the standards indicated:

IEC 60601-2-32 <https://standards.iteh.ai/catalog/standards/sist/en-61223-3-4-2000/4b17ca445dc/sist-en-61223-3-4-2002> NOTE: Harmonized as EN 60601-2-32:1994 (not modified).

Annex ZA (normative)**Normative references to international publications
with their corresponding European publications**

This European Standard incorporates by dated or undated reference, provisions from other publications. These normative references are cited at the appropriate places in the text and the publications are listed hereafter. For dated references, subsequent amendments to or revisions of any of these publications apply to this European Standard only when incorporated in it by amendment or revision. For undated references the latest edition of the publication referred to applies (including amendments).

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

<u>Publication</u>	<u>Year</u>	<u>Title</u>	<u>EN/HD</u>	<u>Year</u>
IEC 60336	1993	X-ray tube assemblies for medical diagnosis - Characteristics of focal spots	EN 60336	1995
IEC 60417-1	1998	Graphical symbols for use on equipment Part 1: Overview and application	EN 60417-1	1999
IEC 60417-2	1998	Part 2: Symbol originals	EN 60417-2	1999
IEC 60522	1999	Determination of the permanent filtration of X-ray tube assemblies	EN 60522	1999
IEC 60601-1	1988	Medical electrical equipment Part 1: General requirements for safety	EN 60601-1 + corr. July + A13	1990 1994 1996
		NOTE: Amendments A11 and A12 are superseded by EN 60601-1/A2:1995.		
IEC 60601-2-28	1993	Part 2: Particular requirements for the safety of X-ray source assemblies and X-ray tube assemblies for medical diagnosis	EN 60601-2-28	1993
IEC 60788	1984	Medical radiology - Terminology	HD 501 S1	1988
IEC 60878	1988	Graphical symbols on electrical equipment in medical practice	-	-
IEC 61223-1	1993	Evaluation and routine testing in medical imaging departments Part 1: General aspects	-	-
IEC 61267	1994	Medical diagnostic X-ray equipment - Radiation conditions for use in the determination of characteristics	EN 61267	1994
ISO 2092	1981	Light metals and their alloys - Code of designation based on chemical symbols	-	-

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First edition
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Evaluation and routine testing in medical imaging departments –

Part 3-4:

Acceptance tests –

Imaging performance of dental X-ray equipment (standards.iteh.ai)

*Essais d'évaluation et de routine dans les services
d'imagerie médicale –*
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Partie 3-4:

Essais d'acceptation –

Performance d'imagerie des appareils de radiographie dentaire

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CONTENTS

	Page
FOREWORD	4
INTRODUCTION	6
Clause	
1 Scope and object	7
1.1 Scope	7
1.2 Object	7
2 Normative references	8
3 Terminology	8
3.1 Degree of requirements	8
3.2 Use of terms	9
3.3 Defined terms	9
4 General aspects of ACCEPTANCE TESTS	9
4.1 General conditions to be considered in test procedures	9
4.2 Documents and data for the tests	10
4.3 Test conditions	10
4.4 Scope of tests	11
4.5 Test equipment including PHANTOMS and TEST DEVICES	11
4.6 Evaluating the test results	13
5 Test methods for dental X-RAY EQUIPMENT with intra-oral X-RAY IMAGE RECEPTOR	13
5.1 Visual and functional tests	13
5.2 *X-RAY TUBE VOLTAGE	13
5.3 *TOTAL FILTRATION	14
5.4 *FOCAL SPOT of the X-RAY TUBE	14
5.5 Limitation and alignment of the X-RAY BEAM	15
5.6 FOCAL SPOT TO SKIN DISTANCE	15
5.7 *Reproducibility of RADIATION OUTPUT	16
5.8 LINE PAIR RESOLUTION	16
5.9 LOW CONTRAST RESOLUTION	16
6 Test methods for dental panoramic X-RAY EQUIPMENT with extra-oral X-RAY IMAGE RECEPTOR	17
6.1 Visual and functional tests	17
6.2 X-RAY TUBE VOLTAGE	17
6.3 TOTAL FILTRATION	17
6.4 FOCAL SPOT of the X-RAY TUBE	17
6.5 Limitation and alignment of the X-RAY BEAM	17
6.6 FOCAL SPOT TO SKIN DISTANCE	18
6.7 Reproducibility of RADIATION OUTPUT	18
6.8 LINE PAIR RESOLUTION	19
6.9 LOW CONTRAST RESOLUTION	19
6.10 RADIOGRAPHIC FILM cassettes with INTENSIFYING SCREENS	19
6.11 Image homogeneity	19
6.12 Indicators for patients' positioning	20
6.13 Panoramic layer	20

Clause	Page
7 Test methods for dental cephalometric X-RAY EQUIPMENT with extra-oral X-RAY IMAGE RECEPTOR.....	20
7.1 Visual and functional tests.....	20
7.2 X-RAY TUBE VOLTAGE.....	20
7.3 TOTAL FILTRATION.....	20
7.4 FOCAL SPOT of the X-RAY TUBE.....	20
7.5 Limitation and alignment of the X-RAY BEAM.....	20
7.6 FOCAL SPOT TO SKIN DISTANCE.....	21
7.7 Reproducibility of RADIATION OUTPUT.....	21
7.8 LINE PAIR RESOLUTION.....	21
7.9 LOW CONTRAST RESOLUTION.....	21
7.10 RADIOGRAPHIC FILM cassettes with INTENSIFYING SCREENS.....	21
8 Test report and statement of compliance.....	21
Annex A (normative) Terminology – Index of defined terms.....	30
Annex B (informative) Examples of requirements (accuracy, tolerances, discrepancies) according to actual IEC standards or state of the art.....	32
Annex C (informative) ACCEPTANCE TEST for processing of non-screen dental X-ray films (non-screen film).....	34
Bibliography.....	35
Figure 1 – Dental X-RAY EQUIPMENT with intra-oral X-RAY IMAGE RECEPTOR measuring arrangement for AIR KERMA and resolution.....	23
Figure 2 – Dental X-RAY EQUIPMENT with intra-oral X-RAY IMAGE RECEPTOR measuring arrangement for AIR KERMA and resolution.....	24
Figure 3 – Dental panoramic X-RAY EQUIPMENT with extra-oral X-RAY IMAGE RECEPTOR measuring arrangement for AIR KERMA and resolution.....	25
Figure 4 – Example of a dental panoramic X-RAY EQUIPMENT with extra-oral digital X-RAY IMAGE RECEPTOR measuring arrangement for AIR KERMA; resolution, image homogeneity and panoramic layer.....	26
Figure 5 – Example of a cephalometric X-RAY EQUIPMENT with extra-oral X-RAY IMAGE RECEPTOR measuring arrangement for AIR KERMA and resolution.....	27
Figure 6 – Dental PHANTOM (example).....	28
Figure 7 – Dental PHANTOM for digital image acquisition or processing parts (example).....	29
Table B.1 – Typical values of FOCAL SPOT dimensions for NOMINAL FOCAL SPOT VALUES.....	32

INTERNATIONAL ELECTROTECHNICAL COMMISSION

EVALUATION AND ROUTINE TESTING
IN MEDICAL IMAGING DEPARTMENTS –Part 3-4: Acceptance tests –
Imaging performance of dental X-ray equipment

FOREWORD

- 1) The IEC (International Electrotechnical Commission) is a worldwide organization for standardization comprising all national electrotechnical committees (IEC National Committees). The object of the IEC is to promote international co-operation on all questions concerning standardization in the electrical and electronic fields. To this end and in addition to other activities, the IEC publishes International Standards. Their preparation is entrusted to technical committees; any IEC National Committee interested in the subject dealt with may participate in this preparatory work. International, governmental and non-governmental organizations liaising with the IEC also participate in this preparation. The IEC collaborates closely with the International Organization for Standardization (ISO) in accordance with conditions determined by agreement between the two organizations.
- 2) The formal decisions or agreements of the IEC on technical matters express, as nearly as possible, an international consensus of opinion on the relevant subjects since each technical committee has representation from all interested National Committees.
- 3) The documents produced have the form of recommendations for international use and are published in the form of standards, technical specifications, technical reports or guides and they are accepted by the National Committees in that sense.
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International Standard IEC 61223-3-4 has been prepared by subcommittee 62B: Diagnostic imaging equipment, of IEC technical committee 62: Electrical equipment in medical practice.

The text of this standard is based on the following documents:

FDIS	Report on voting
62B/393/FDIS	62B/402/RVD

Full information on the voting for the approval of this standard can be found in the report on voting indicated in the above table.

This publication has been drafted in accordance with the ISO/IEC Directives, Part 3.

Annex A forms an integral part of this standard.

Annexes B and C are for information only.

In this standard, the following print types are used:

- requirements, compliance with which can be tested, and definitions: roman type;
- explanations, advice, notes, general statements, exceptions and references: smaller type;
- *test specifications: italic type;*
- TERMS DEFINED IN IEC 60601-1, IN IEC 60788, IN IEC 61223-1 OR IN OTHER IEC PUBLICATIONS REFERENCED IN ANNEX A: SMALL CAPITALS.

The committee has decided that the contents of this publication will remain unchanged until 2005. At this date, the publication will be

- reconfirmed;
- withdrawn;
- replaced by a revised edition, or
- amended.

A bilingual version of this standard may be issued at a later date.

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INTRODUCTION

This part of IEC 61223 is part of a series of International Standards which gives methods of acceptance testing and constancy testing for subsystems and systems (for example, diagnostic X-RAY EQUIPMENT) including film processing.

Some provisions or statements in this standard require additional information. Such information is presented in annex B. An asterisk in the left margin of a clause or subclause indicates the presence of such additional information.

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EVALUATION AND ROUTINE TESTING IN MEDICAL IMAGING DEPARTMENTS –

Part 3-4: Acceptance tests – Imaging performance of dental X-ray equipment

1 Scope and object

1.1 Scope

This part of IEC 61223 applies to those components of dental X-RAY EQUIPMENT using radiographic imaging systems which influence the image quality and PATIENT dose.

This standard applies to the performance of the ACCEPTANCE TEST on dental X-RAY EQUIPMENT with intra-oral X-RAY IMAGE RECEPTOR and dental X-RAY EQUIPMENT with extra-oral X-RAY IMAGE RECEPTOR (for example, dental panoramic X-RAY EQUIPMENT or cephalometric X-RAY).

This standard applies to dental film and digital image acquisition and processing.

1.2 Object

This standard defines

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- a) the essential parameters which describe the performance of the above-mentioned dental X-RAY EQUIPMENT with regard to imaging properties and PATIENT dose;
- b) methods of testing and whether measured quantities related to those parameters comply with the specified tolerances.

These methods rely mainly on non-invasive measurements, using appropriate test equipment, performed during or after the installation is completed. Signed statements covering steps in the installation procedure may be used as part of the acceptance testing.

The aim is to verify compliance of the installation with specifications affecting the image quality and PATIENT dose, and to detect malfunctions that are not in agreement with those specifications.

This standard does not in itself specify tolerances for the parameters under investigation. Neither is it intended to consider

- c) aspects of mechanical and electrical safety;
- d) aspects of mechanical, electrical and software performance, unless they are essential to the performance of the tests directly affecting image quality and PATIENT dose.