SLOVENSKI STANDARD
SIST EN 61262-6:1995

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Medical electrical equipment - Characteristics of electro optical X ray image intensifiers - Part 6: Determination of the contrast ratio and veiling glare index (IEC 1262-6:1994)

Medical electrical equipment - Characteristics of electro-optical X-ray image intensifiers -- Part 6: Determination of the contrast ratio and veiling glare index

Medizinische elektrische Geräte - Merkmale von elektronenoptischen Röntgenbildverstärkern -- Teil 6: Bestimmung des Kontrastverhältnisses und Untergrundkoeffizienten

Appareils électromédicaux - Caractéristiques des intensificateurs électro-optiques d'image radiologique -- Partie 6: Détermination du rapport de contraste et du voile lumineux

Ta slovenski standard je istoveten z: **EN 61262-6:1994**

ICS:

11.040.50	Radiografska oprema	Radiographic equipment

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Medical electrical equipment - Characteristics of electro-optical X-ray image intensifiers Part 6: Determination of the contrast ratio and veiling glare index (IEC 1262-6:1994)

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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(CO)117, as prepared by Sub-Committee 62B: Diagnostic imaging equipment, of IEC Technical Committee 62: Electrical equipment in medical practice was submitted to the IEC-CENELEC parallel vote in January 1994.

The reference document was approved by CENELEC as EN 61262-6 on 5 July 1994.

The following dates were fixed:

- latest date of publication of an identical national standard	(dop)	1995-07-01
- latest date of withdrawal of conflicting national standards	(dow)	1995-07-01

For products which have complied with the relevant national standard before 1995-07-01 as shown by the manufacturer or by a certification body, this previous standard may continue to apply for production until 2000-07-01.

Annexes designated "normative" are part of the body of the standard. Annexes designated "informative" are given only for information. In this standard, annexes A and B are informative and annex ZA is normative.

ENDORSEMENT NOTICE

The text of the International Standard IEC 1262-6:1994 was approved by CENELEC as a European Standard without any modification.

ANNEX ZA (normative)

OTHER INTERNATIONAL PUBLICATIONS QUOTED IN THIS STANDARD WITH THE REFERENCES OF THE RELEVANT 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.

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

IEC Publication	Date	Title	EN/HD	Date
788	1984	Medical radiology - Terminology	HD 501 S1	1988

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INTERNATIONAL ELECTROTECHNICAL COMMISSION

MEDICAL ELECTRICAL EQUIPMENT –

CHARACTERISTICS OF ELECTRO-OPTICAL X-RAY IMAGE INTENSIFIERS –

Part 6: Determination of the contrast ratio and veiling glare index

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 cooperation on all questions concerning standardization in the electrical and electronic field. 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.

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3) They have the form of recommendations for international use published in the form of standards, technical reports or guides and they are accepted by the National Committees in that sense.

4) In order to promote international unification, IEC National Committees undertake to apply IEC International Standards transparently to the maximum extent possible in their national and regional standards. Any divergence between the IEC Standard and the corresponding national or regional standard shall be clearly indicated in the latter.

International Standard IEC 1262-6 has been prepared by sub-committee 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:

DIS	Report on voting
62B(CO)117	62B(CO)129

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

Annexes A and B are for information only.

In this standard, the following print types are used:

– Requirements, compliance with which can be tested, and definitions: in roman type.

– Explanations, advice, introductions, general statements, and exceptions: in smaller type.

– *Test specifications: in italic type.*

– TERMS USED THROUGHOUT THIS STANDARD WHICH HAVE BEEN DEFINED IN 3.1 AND IN ANNEX A: SMALL CAPITALS.