



National Standards Authority of Ireland

IRISH STANDARD

I.S. EN 61262-2:1995

ICS 11.040.50

**MEDICAL ELECTRICAL EQUIPMENT –  
CHARACTERISTICS OF ELECTRO-OPTICAL  
X-RAY IMAGE INTENSIFIERS. PART 2:  
DETERMINATION OF THE CONVERSION  
FACTOR (IEC 1262-2:1994)**

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Údarás um Chaighdeán Náisiúnta na hÉireann



## DECLARATION

OF

SPECIFICATION

ENTITLED

MEDICAL ELECTRICAL EQUIPMENT – CHARACTERISTICS OF ELECTRO-OPTICAL  
X-RAY IMAGE INTENSIFIERS. PART 2: DETERMINATION OF THE CONVERSION  
FACTOR (IEC 1262-2:1994)

AS

THE IRISH STANDARD SPECIFICATION FOR

MEDICAL ELECTRICAL EQUIPMENT – CHARACTERISTICS OF ELECTRO-OPTICAL  
X-RAY IMAGE INTENSIFIERS. PART 2: DETERMINATION OF THE CONVERSION  
FACTOR (IEC 1262-2:1994)

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Forfás in exercise of the power conferred by section 20 (5) of the Industrial Research and Standards Act, 1961 ( No. 20 of 1961 ) and the Industrial Development Act, 1993 (No. 19 of 1993), and with the consent of the Minister for Enterprise and Employment, hereby declares as follows:

1. This instrument may be cited as the Standard Specification (Medical Electrical Equipment – Characteristics of Electro-Optical X-Ray Image Intensifiers. Part 2: Determination of the Conversion Factor (IEC 1262-2:1994)) Declaration, 1995.

2. (1) The Specification set forth in the Schedule to this declaration is hereby declared to be the standard specification for Medical Electrical Equipment – Characteristics of Electro-Optical X-Ray Image Intensifiers. Part 2: Determination of the Conversion Factor (IEC 1262-2:1994)

(2) The said standard specification may be cited as Irish Standard/EN 61262-2:1995 or as I.S./EN 61262-2:1995.

3. (1) The Standard Specification (Measurement of the conversion factor of electro-optical X-ray intensifiers, Edition 1) Declaration, 1990 is hereby revoked.

(2) Reference in any other standard specification to the Instrument hereby revoked and to Irish Standard/HD 512 S1:1989 thereby prescribed, shall be construed, respectively, as references to this Instrument and to Irish Standard/EN 61262-2:1995.



EUROPEAN STANDARD

EN 61262-2

NORME EUROPEENNE

EUROPÄISCHE NORM

September 1994

ICS 11.040.50

Supersedes HD 512 S1:1989

Descriptors: Medical electrical equipment, image intensifier, X-ray,  
conversion factor

ENGLISH VERSION

Medical electrical equipment - Characteristics of  
electro-optical X-ray image intensifiers  
Part 2: Determination of the conversion factor  
(IEC 1262-2:1994)

Appareils électromédicaux  
Caractéristiques des  
intensificateurs  
électro-optiques d'image  
radiologique  
Partie 2: Détermination du  
facteur de conversion  
(CEI 1262-2:1994)

Medizinische elektrische  
Geräte - Merkmale von  
elektronenoptischen  
Röntgenbildverstärkern  
Teil 2: Bestimmung des  
Konversionsfaktors  
(IEC 1262-2:1994)

This European Standard was approved by CENELEC on 1994-07-05.  
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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 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,  
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(CO)113, 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 61262-2 on 5 July 1994.

This European Standard replaces HD 512 S1:1989.

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 HD 512 S1:1989 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 1262-2:1994 was approved by CENELEC as a European Standard without any modification.

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