SLOVENSKI STANDARD

SIST EN 60601-2-33:1998/A11:1998

prva izdaja
september 1998

Medical electrical equipment - Part 2: Particular requirements for the safety of magnetic resonance equipment for medical diagnosis - Amendment A11 - Amendment to annex AA of EN

ICS 11.040.50

Referenčna številka
SIST EN 60601-2-33:1998/A11:1998(en)

EUROPEAN STANDARD

NORME EUROPÉENNE

EUROPÄISCHE NORM

EN 60601-2-33/A11

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Descriptors: Medical electrical equipment, magnetic resonance equipment, magnetic resonance examination, safety requirements, protection against electric shock, protection against mechanical hazard, radiation protection, fire protection, environmental conditions

English version

Medical electrical equipment
Part 2: Particular requirements for the safety of magnetic resonance equipment for medical diagnosis

Appareils électromédicaux
Partie 2: Règles particulières de sécurité relatives aux appareils à résonance magnétique pour diagnostic médical

Medizinische elektrische Geräte
Teil 2: Besondere Festlegungen für die Sicherheit von medizinischen diagnostischen Magnetresonanzgeräten

This amendment A11 modifies the European Standard EN 60601-2-33:1995; it was approved by CENELEC on 1997-03-11. CENELEC members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this amendment 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 amendment 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

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Foreword

This amendment was prepared by the Technical Committee CENELEC TC 62, Electrical equipment in medical practice.

The text of the draft was submitted to the formal vote and was approved by CENELEC as amendment A11 to EN 60601-2-33:1995 on 1997-03-11.

The following dates were fixed:

- latest date by which the amendment has to be implemented at national level by publication of an identical national standard or by endorsement (dop) 1998-03-01
- latest date by which the national standards conflicting with the amendment have to be withdrawn (dow) 1998-06-13

Replace annex AA by the following new annex:

Annex AA
(informative)

Examples of warning signs and prohibitive signs in line with the European Council Directive 92/58/EEC

	Warning sign: Strong magnetic field
	Warning sign: Non-ionizing radiation
	Prohibitive sign: No access for persons with pacemakers
	Prohibitive sign: No access for persons with metal implants
	Prohibitive sign: No metallic articles or watches