

STN	Zdravotnícke elektrické prístroje Časť 1-8: Všeobecné požiadavky na základnú bezpečnosť a nevyhnutné prevádzkové vlastnosti Pridružená norma: Všeobecné požiadavky, skúšky a pokyny pre poplachové systémy v zdravotníckych elektrických prístrojoch a zdravotníckych elektrických systémoch Zmena A11	STN EN 60601-1-8/A11 36 4800
------------	---	--

Medical electrical equipment. Part 1-8: General requirements for basic safety and essential performance. Collateral Standard: General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems

Táto norma obsahuje anglickú verziu európskej normy.
This standard includes the English version of the European Standard.

Táto norma bola oznámená vo Vestníku ÚNMS SR č. 10/17

STN EN 60601-1-8 z apríla 2008 sa bez zmeny A11 môže používať do 7. 1. 2020.

Obsahuje: EN 60601-1-8:2007/A11:2017

125552

Úrad pre normalizáciu, metrológiu a skúšobníctvo Slovenskej republiky, 2017
Podľa zákona č. 264/1999 Z. z. o technických požiadavkách na výrobky a o posudzovaní zhody a o zmene a doplnení niektorých zákonov v znení neskorších predpisov sa slovenská technická norma a časti slovenskej technickej normy môžu rozmnožovať alebo rozširovať len so súhlasom slovenského národného normalizačného orgánu.

ICS 11.040.01

English Version

Medical electrical equipment -
Part 1-8: General requirements for basic safety and essential
performance - Collateral Standard: General requirements, tests
and guidance for alarm systems in medical electrical equipment
and medical electrical systems

Appareils électromédicaux -
Partie 1-8: Exigences générales pour la sécurité de base et
les performances essentielles - Norme collatérale:
Exigences générales, essais et guide pour les systèmes
d'alarme des appareils et des systèmes électromédicaux

Medizinische elektrische Geräte -
Teil 1-8: Allgemeine Festlegungen für die Sicherheit
einschließlich der wesentlichen Leistungsmerkmale -
Ergänzungsnorm: Alarmsysteme - Allgemeine
Festlegungen, Prüfungen und Richtlinien für Alarmsysteme
in medizinischen elektrischen Geräten und in medizinischen
elektrischen Systemen

This amendment A11 modifies the European Standard EN 60601-1-8:2007; it was approved by CENELEC on 2017-01-07. 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 CEN-CENELEC Management Centre 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 CEN-CENELEC Management Centre has the same status as the official versions.

CENELEC members are the national electrotechnical committees of Austria, Belgium, Bulgaria, Croatia, Cyprus, the Czech Republic, Denmark, Estonia, Finland, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, the Netherlands, Norway, Poland, Portugal, Romania, Serbia, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and the United Kingdom.



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

EN 60601-1-8:2007/A11:2017**European foreword**

This document (EN 60601-1-8:2007/A11:2017) has been prepared by CLC/TC 62 "Electrical equipment in medical practice".

The following dates are fixed:

- latest date by which this document has to be implemented at national level by publication of an identical national standard or by endorsement (dop) 2018-07-01
- latest date by which the national standards conflicting with this document have to be withdrawn (dow) 2020-01-07

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. CENELEC [and/or CEN] shall not be held responsible for identifying any or all such patent rights.

This document has been prepared under a mandate given to CENELEC by the European Commission and the European Free Trade Association, and supports essential requirements of EU Directive.

For the relationship with EU Directive see informative Annex ZZ, which is an integral part of this document.

koniec náhľadu – text ďalej pokračuje v platenej verzii STN