

ציוד חשמלי לשימוש רפואי: דרישות מיוחדות לבטיחות בסיסית ולביצועים חיוניים של ציוד ריפוי הפועל בגלי-מיקרו

Medical electrical equipment: Particular requirements for the basic safety and essential performance of microwave therapy equipment

נסמך לה הוא הצעה בלבד

גיליון תיקון זה הוכן ואושר על ידי הוועדה הטכנית 5801 – ציוד חשמלי לשימוש רפואי, בהרכב זה:

- איגוד לשכות המסחר - אייל גרינבאום, אורנה שרון
- המועצה הישראלית לצרכנות - מיכאל שיזף
- מינוי אישי - ולנטין ויינטראוב, יוסי פרי פז
- מינוי אישי – מעבדה - אירנה אנטונוב
- משרד הבריאות - אלכסנדר וילנסקי (יו"ר)
- משרד הכלכלה והתעשייה - שלומי אביסרור

כמו כן תרם להכנת הגיליון התיקון דני אבן-חן.

זיוה שלו ריכזה את עבודת הכנת גיליון התיקון.

פרסומים

הודעה על גיליון תיקון

גיליון תיקון זה, למעט השינויים והתוספות הלאומיים המצוינים בו, זהה לגיליון התיקון של הנציבות הבין-לאומית לאלקטרוטכניקה

IEC 60601-2-6:2012/AMD2: 2022-09

גיליון תיקון זה מעדכן את

התקן הישראלי ת"י 60601 חלק 2.6 מאוקטובר 2018

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הקדמה לגיליון התיקון הישראלי

גיליון תיקון ישראלי זה הוא גיליון התיקון של הנציבות הבין-לאומית לאלקטרוטכניקה IEC 60601-2-6:2012/AMD2 מספטמבר 2022, שאושר כגיליון תיקון ישראלי בשינויים ובתוספות לאומיים. גיליון התיקון של הנציבות הבין-לאומית לאלקטרוטכניקה מובא להלן בשפה האנגלית.

שינויים לחלק העברי של התקן:

בסוף החלק העברי של התקן יוסף:

פירוט השינויים והתוספות הלאומיים לסעיפי התקן הבין-לאומי

201.2 Normative references

במקום אחד התקנים הבין-לאומיים המאוזכרים בתקן והמפורט בסעיף זה חל תקן ישראלי, כמפורט להלן:

התקן הישראלי החל במקומו	התקן הבין-לאומי המאוזכר
ת"י 60601 חלק 1 – ציוד חשמלי לשימוש רפואי: דרישות כלליות לבטיחות בסיסית ולביצועים חיוניים	IEC 60601-1:2005 IEC 60601-1:2005/AMD1:2012 IEC 60601-1:2005/AMD2:2020

INTERNATIONAL STANDARD

NORME INTERNATIONALE

AMENDMENT 2
AMENDEMENT 2

**Medical electrical equipment –
Part 2-6: Particular requirements for the basic safety and essential performance
of microwave therapy equipment**

**Appareils électromédicaux –
Partie 2-6: Exigences particulières pour la sécurité de base et les performances
essentielles des appareils de thérapie à micro-ondes**





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INTERNATIONAL STANDARD

NORME INTERNATIONALE

AMENDMENT 2
AMENDEMENT 2

**Medical electrical equipment –
Part 2-6: Particular requirements for the basic safety and essential performance
of microwave therapy equipment**

**Appareils électromédicaux –
Partie 2-6: Exigences particulières pour la sécurité de base et les performances
essentiels des appareils de thérapie à micro-ondes**

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Amendment 2 to IEC 60601-2-6:2012 has been prepared by subcommittee 62D: Electromedical equipment, of IEC technical committee 62: Electrical equipment in medical practice.

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

Draft	Report on voting
62D/1847/CDV	62D/1960/RVC

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

The language used for the development of this Amendment is English.

This document was drafted in accordance with ISO/IEC Directives, Part 2, and developed in accordance with ISO/IEC Directives, Part 1 and ISO/IEC Directives, IEC Supplement, available at www.iec.ch/members_experts/refdocs. The main document types developed by IEC are described in greater detail at www.iec.ch/standardsdev/publications/.

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INTRODUCTION to Amendment 2

At the October 2019 meeting of IEC SC 62D in Shanghai, China, the subcommittee discussed the need for administrative/technical changes to most 62D standards after completion of the amendment projects within the IEC 60601-1 series. Those projects were all completed and the amendments published in 2020.

The full list of IEC SC 62D documents that will be amended or revised may be found within the IEC document 62D/1792/DC. The results and comments on the DC may be found within 62D/1808/INF. The review report for this amendment is 62D/1830/RR.

201.1 Scope, object and related standards

Replace the existing text of footnote 1, modified by Amendment 1, with the following:

The general standard is IEC 60601-1:2005, IEC 60601-1:2005/AMD1:2012 and IEC 60601-1:2005/AMD2:2020, *Medical electrical equipment – Part 1: General requirements for basic safety and essential performance*.

201.1.3 Collateral standards

Replace, in the second paragraph, the second sentence with the following:

IEC 60601-1-3 and IEC 60601-1-12 do not apply.

201.2 Normative references

Replace the existing text with the following:

Clause 2 of the general standard applies, except as follows:

Addition:

IEC 60601-1:2005, *Medical electrical equipment – Part 1: General requirements for basic safety and essential performance*

IEC 60601-1:2005/AMD1:2012

IEC 60601-1:2005/AMD2:2020

IEC TR 60878:2015, *Graphical symbols for electrical equipment in medical practice*

201.3 Terms and definitions

Replace, in the first sentence, modified by Amendment 1, “IEC 60601-1” with “IEC 60601-1:2005, IEC 60601-1:2005/AMD1:2012 and IEC 60601-1:2005/AMD2:2020”.

Index of defined terms used in this particular standard

Replace the following entries, modified by Amendment 1, as follows:

ACCESS COVER	IEC 60601-1:2005, 3.1
APPLIED PART	IEC 60601-1:2005, 3.8
BASIC SAFETY	IEC 60601-1:2005, 3.10
CATEGORY AP	IEC 60601-1:2005, 3.11
CATEGORY APG	IEC 60601-1:2005, 3.12
ESSENTIAL PERFORMANCE	IEC 60601-1:2005 and IEC 60601-1:2005/AMD1:2012, 3.27
HAZARD.....	IEC 60601-1:2005 and IEC 60601-1:2005/AMD2:2020, 3.39
LEAKAGE CURRENT	IEC 60601-1:2005, 3.47
MANUFACTURER.....	IEC 60601-1:2005 and IEC 60601-1:2005/AMD2:2020, 3.55
MEDICAL ELECTRICAL EQUIPMENT (ME EQUIPMENT).....	IEC 60601-1:2005, 3.63
MEDICAL ELECTRICAL SYSTEM (ME SYSTEM).....	IEC 60601-1:2005, 3.64
OPERATOR	IEC 60601-1:2005, 3.73
PATIENT	IEC 60601-1:2005 and IEC 60601-1:2005/AMD1:2012, 3.76
PATIENT AUXILIARY CURRENT	IEC 60601-1:2005, 3.77
TOOL	IEC 60601-1:2005, 3.127

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