

## ***Iec 60601 2 33 Ed 21 B2006 Medical Electrical Equipment Part 2 33 Particular Requirements For The Safety Of Magnetic Resonance Equipment For Medical Diagnosis***

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***This edition of IEC 60601-2-33 is based on the second amendment to Edition 2. It has also been adapted to the third edition of 60601-1 (2005) IEC, with technical modifications being introduced where appropriate. This publication has been drafted in accordance with the ISO/IEC Directives, Part 2.***

***IEC 60601 - Wikipedia***

***– 4 – 60601-2-33 Amend. 1 IEC:2005(E) Page 45 Annex BB – Guidance and rationale for particular subclauses Add, on page 57, a rationale for subclause item 6.8.2 ss) as follows: Concerning 6.8.2 ss) In addition to the information given in Clause 6.8.2 cc) on emergency medical procedures and***

***IEC 60601-2-33 Ed. 2.0 en:2002 - Techstreet***

***IEC 60601-2-33 - Edition 2 TESTING AND MEASURING EQUIPMENT/ALLOWED***

**SUBCONTRACTING R=Required by Lab; ; W= Not required in CBTL, can be witnessed in MTL situations S=May be subcontracted Clause Measurement/testing Testing / measuring equipment / material needed Subcontracting 26 Sound level sound level meter to IEC 60651 and/or IEC 60804.**

### ***IEC 60601-2-33 Review – RITE Advantage***

***This third edition cancels and replaces the second edition published in 2002, its Amendment 1 (2005) and Amendment 2 (2007) and constitutes a technical revision. It has also been adapted to the third edition of IEC 60601-1 (2005), with technical modifications being introduced where appropriate.***

### ***IEC 60601-2-33:2010/AMD2:2015***

***This third edition of IEC 60601-2-33 is based on the second amendment to Edition 2. It has also been adapted to the third edition of IEC 60601-1 (2005), with technical modifications being introduced where appropriate. The contents of the corrigenda of March 2012 and February 2016 have been included in this copy.***

### ***IEC 60601-2-33:2010+AMD1:2013 CSV***

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***IEC 60601-2-33 Ed. 3.0 b:2010 - Medical electrical ...***

***IEC 60601-2-33, 3.2 Edition, June 2015 - Medical electrical equipment – Part 2-33:  
Particular requirements for the basic safety and essential performance of magnetic  
resonance equipment for medical diagnosis This International Standard applies to the  
BASIC SAFETY and ESSENTIAL PERFORMANCE of MR EQUIPMENT and MR SYSTEMS,  
hereafter referred to also as ME EQUIPMENT.***

***IEC Standard - Home***

***Particular standards (numbered 60601-2-X) define the requirements for specific  
products or specific measurements built into products, e.g. MR scanners (IEC  
60601-2-33) or Electroencephalograms (IEC 60601-2-26). Collaterals and Particulars may  
have their own revisions which are different from the General Standard.***

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***INTERNATIONAL IEC STANDARD 60601-2-33***

***IEC 60601-2-33 Ed. 3.0 b:2010 Medical electrical equipment - Part 2-33: Particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis "IEC 60601-2-33:2010 establishes particular basic safety and essential performance requirements for magnetic resonance equipment to provide protection for the patient and the magnetic resonance worker.***

### ***Recognized Consensus Standards***

***IEC 60601-2-33:2010+A1:2013+A2:2015 establishes particular basic safety and essential performance requirements for magnetic resonance equipment to provide protection for the patient and the magnetic resonance worker. This third edition of IEC 60601-2-33 is based on the second amendment to Edition 2.***

### ***IEC 60601-2-33:2010 | IEC Webstore***

***International Standard IEC 60601-2-33 has been prepared by subcommittee 62B: Diagnostic imaging equipment, of IEC technical committee 62: Electrical equipment in medical practice. This second edition cancels and replaces the first edition published in 1995 and constitutes a technical revision.***

### ***IEC 60601-2-33 Ed. 3.1 b:2013***

***IEC: 60601-2-33 Ed. 3.2 b:2015: Medical electrical equipment - Part 2-33: Particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis: 12/23/2019: Radiology: 12-296: IEC: 60601-2-54***

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**IEC 60601-2-33 - Edition 2 TESTING AND MEASURING EQUIPMENT ...**

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**information. This audio course was built from reviewing each line of the document and summarizing it into a 1 hour course.**

**IEC 60601-2-33:2015 - Estonian Centre for Standardisation**

**IEC 60447 zavedena v ?SN EN 60447 ed. 2 (33 0173) Základní a bezpečnostní zásady pro rozhraní ?lov?k-stroj, zna?ení a identifikaci - Zásady pro ovládání. IEC 60529:1989 zavedena v ?SN EN 60529:1993 (33 0330) Stupn? ochrany krytem (krytí - IP kód). IEC 60601-1-2 zavedena v ?SN EN 60601-1-2 (36 4800) Zdravotnické elektrické p?ístroje - ?ást 1-2: Všeobecné požadavky ...**

**IEC 60601-2-33 Ed. 3.2 b:2015**

**IEC 60601-2-33 Ed. 2.0 en:2002 Medical electrical equipment - Part 2-33: Particular requirements for the safety of magnetic resonance equipment for medical diagnosis . standard by International Electrotechnical Commission, 05/22/2002 Amendments Available. View all product details**

**Edition 3.2 2015-06 FINAL VERSION VERSION FINALE**

**This Particular Standard applies to MR EQUIPMENT as defined in 2.2.101 and MR SYSTEMS as defined in 2.2.102. This Standard does not cover the application of MR EQUIPMENT beyond the INTENDED USE. IEC 60601-2-33**

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