

## 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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Compliance with Medical Standards IEC 62304, ISO 14971, IEC 60601, FDA Title 21 CFR Part 11 [IEC 60601-2-33 Ed](#)

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.

~~INTERNATIONAL IEC STANDARD 60601-2-33~~

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 | IEC Webstore~~

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 60601-2-33 : Medical electrical equipment – Part 2-33 ...~~

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IEC 60601-2-33:2010/AMD1:2013 Standard | Amendment 1 - Medical electrical equipment - Part 2-33: Particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis

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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.

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publication has been drafted in accordance with the ISO/IEC Directives, Part 2.

## ~~Edition 3.2 2015-06 CONSOLIDATED VERSION CONSOLIDÉE~~

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. 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 corrigendum of March 2012 ...

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## ~~IEC Standard - Home~~

The general standard IEC 60601-1 - Medical electrical equipment - Part 1: General requirements for basic safety and essential performance - gives general requirements of the series of standards. 60601 is a widely accepted benchmark for medical electrical equipment and compliance with IEC 60601-1 has become a requirement for the commercialisation of electrical medical equipment in many countries.

## ~~IEC 60601 - Wikipedia~~

"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.

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IEC 60601-2-31 Edition 3.0 2020-01 INTERNATIONAL STANDARD NORME INTERNATIONALE Medical electrical equipment – Part 2-31: Particular requirements for the basic safety and essential performance of external cardiac pacemakers with internal power source . Appareils électromédicaux – Partie 2-31: Exigences particulières pour la sécurité de base et les performances essentielles des ...

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## ~~Edition 3.2 2015-06 FINAL VERSION VERSION FINALE~~

IEC 60601-1-2 Edition 4.1 2020-09 CONSOLIDATED VERSION REDLINE VERSION Medical electrical equipment – Part 1-2: General requirements for basic safety and essential performance – Collateral Standard: Electromagnetic disturbances – Requirements and tests . IEC 60601-1-2:2014-02+AMD1:2020-09 CSV(en) ® colour inside This is a preview - click here to buy the full publication – 2 – IEC ...

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## ~~IEC 60601-2-23 Review - RITE Advantage~~

Full Description IEC 60601-2-23:2011 applies to the basic safety and essential performance of transcutaneous partial pressure monitoring equipment. It applies to transcutaneous monitors used with adults, children and neonates, and it includes the use of these devices in foetal monitoring during birth.

## ~~IEC 60601-2-23 Ed. 3.0 b:2011~~

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