

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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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 IEC 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.

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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.

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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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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 60601—Wikipedia~~

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Medical Electrical Equipment Part 2 33

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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