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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.

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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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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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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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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

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equipment in many countries.

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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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General requirements for basic safety and essential performance

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