

ICS 11.040.55;

PN-EN 60601-2-33:2003/AC

czerwiec 2009

Wprowadza
EN 60601-2-33:2002/A2:2008/AC:2008, IDT

Dotyczy

PN-EN 60601-2-33:2003

Medyczne urządzenia elektryczne -- Część 2-33: Szczegółowe wymagania bezpieczeństwa urządzeń rezonansu magnetycznego do diagnostyki medycznej

Na wniosek Komitetu Technicznego nr 67
ds. Elektrycznej Aparatury Medycznej

Poprawka do Normy Europejskiej EN 60601-2-33:2002/A2:2008/AC:2008 Medical electrical equipment - Part 2-33: Particular requirements for the safety of magnetic resonance equipment for medical diagnosis

ma status Poprawki do Polskiej Normy



Corrigendum to EN 60601-2-33:2002/A2:2008

English version

Foreword

Add:

Within the European Union some legal provisions may be different from what is indicated in this standard. As these legal provisions are adopted by the European Parliament and the Council they supersede the rules and recommendations of this standard wherever they are conflicting. In particular EU legislation does not make any difference between a “MR Worker” and a “Worker” and occupational health and safety provisions apply to all categories without distinction. Currently, the Directive 2004/40/EC for the protection of workers exposed to risks arising from electromagnetic fields – albeit under review – foresees more stringent exposure limit values for workers. These occupational limits will have to be implemented by all Member States by 30 April 2012 unless the review process leads to the adoption by the European Parliament and the Council of other values before that date.

November 2008

