

Implants for surgery—Active implantable medical devices

Part 1: General requirements for safety, marking and for information to be provided by the manufacturer



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- Australian Society for Biomaterials
- Medical Technology Association of Australia
- Neurosurgical Society of Australasia
- Royal Australasian College of Surgeons
- Royal Perth Hospital
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AS ISO 14708.1:2015

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PREFACE

This Standard was prepared by the Standards Australia Committee HE-012, Surgical Implants, to supersede AS ISO 14708.1—2003.

The objective of this Standard is to provide general requirements on safety, marking and information to be provided by manufacturers of active implantable medical devices.

This Standard is identical with, and has been reproduced from ISO 14708-1:2014, *Implants for surgery—Active implantable medical devices*, Part 1: General requirements for safety, marking and for information to be provided by the manufacturer.

As this Standard is reproduced from an International Standard, the following applies:

- (a) In the source text 'this part of ISO 14708' should read 'this Australian Standard'.
- (b) A full point substitutes for a comma when referring to a decimal marker.

References to International Standards should be replaced by references to Australian or Australian/New Zealand Standards, as follows:

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11137	Sterilization of health care products— Radiation	11137	Sterilization of health care products— Radiation
11137-1	Part 1: Requirements for development, validation and routine control of a sterilization process for medical devices	11137.1	Part 1: Requirements for development, validation and routine control of a sterilization process for medical devices

Only normative references that have been adopted as Australian or Australian/New Zealand Standard have been listed.

The term 'informative' has been used in this Standard to define the application of the annex to which it applies. An 'informative' annex is only for information and guidance.

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