

Notification of a Body in the framework of a technical harmonization directive

From : PERMANENT
REPRESENTATION OF
IRELAND TO THE EUROPEAN
UNION
1040 BRUXELLES

To : **European Commission**
Enterprise Directorate-General
-
B 1049 Brussels
Other Member States

Reference :

Legislation : 90/385/EEC Active implantable medical devices

Body name, address, telephone, fax, email, website :

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Body :

NB 0050

Created : Unknown (Notifications pre-dating 2006 are not available in these lists) | **Last update :** 07/08/2014

Period of validity of the notification :

Valid until : Unlimited

The body is formally accredited against :

EN 45012 - EN ISO/IEC 17021

EN 45011 - EN ISO/IEC 17065

Name of National Accreditation Body (NAB) :

The accreditation covers the product categories and conformity assessment procedures concerned by this notification : No

Tasks performed by the Body :

Created : Unknown (Notifications pre-dating 2006 are not available in these lists) | Last update :

Product family, product /Intended use/Product range	Procedure/Modules	Annexes or articles of the directives	Limitations
All active implantable medical devices	EC declaration of conformity (complete quality assurance system) EC declaration of conformity to type (assurance of production quality)	Annex 2 Annex 5	