

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

**From :** STAENDIGE VERTRETUNG  
DER BUNDESREPUBLIK  
DEUTSCHLAND BEI DER  
EUROPAEISCHEN 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 :**

TÜV NORD CERT GmbH  
Langemarckstraße 20  
45141 ESSEN  
Germany  
Phone : +49 (0) 201 825-3335  
Fax : +49 (0) 201 825-3290  
Email : info@tuev-nord.de  
Website : www.tuev-nord-cert.de

**Body :**

**NB 0044**

**Created :** Unknown (Notifications pre-dating 2006 are not available in these lists) | **Last update :** 04/06/2013

**Period of validity of the notification :**

Valid until : 31/12/2090

**The body is formally accredited against :**

EN 45010  
EN 45012 - EN ISO/IEC 17021  
EN 45011 - EN ISO/IEC 17065  
EN 45000  
EN 45003

**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 :

<b>Product family, product /Intended use/Product range</b>	<b>Procedure/Modules</b>	<b>Annexes or articles of the directives</b>	<b>Limitations</b>
All active implantable medical devices	EC declaration of conformity (complete quality assurance system) EC declaration of conformity to type (assurance of production quality) EC type-examination EC verification	Annex 2 Annex 5 Annex 3 Annex 4	