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

From : PERMANENTE VERTEGENWOORDIGING VAN HET KONINKRIJK DER NEDERLANDEN BIJ DE EUROPESE UNIE
1160 BRUXELLES

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

Other Member States

Reference : Legislation : 93/42/EEC Medical devices

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

TÜV Rheinland Nederland B.V.
Boogschutterstraat 11A Postbus 541 7300 AM
APELDOORN
Netherlands
Phone : +31/88 888 7 888
Fax : +31/88 888 7 879
Email : info@NL.TUV.com
Website : http://www.TUV.com/NL

Body : **NB 0336**

Created : Unknown (Notifications pre-dating 2006 are not available in these lists) **| Last update :** 03/09/2012

Period of validity of the notification :

Valid until : Unlimited

The body is formally accredited against :

EN 45010

EN 45012 - EN ISO/IEC 17021

EN 45011 - EN ISO/IEC 17065

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 :**

Product family, product /Intended use/Product range	Procedure/Modules	Annexes or articles of the directives	Limitations
All medical devices	Full quality assurance system EC type-examination EC verification Production quality assurance Product quality assurance	Annex II Annex III Annex IV Annex V Annex VI	