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

From : Agence Fédérale des Médicaments et des Produits de Santé (AFMPS) - Federaal Agentschap voor Geneesmiddelen en Gezondheidsproducten (FAGG)
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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 :

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Body : **NB 1639**

Created : Unknown (Notifications pre-dating 2006 are not available in these lists) **| Last update :** 28/09/2009

Period of validity of the notification :

Valid until : 09/05/2011(Expired/Withdrawn)

The body is formally accredited against :

Name of National Accreditation Body (NAB) : BELAC

The accreditation covers the product categories and conformity assessment procedures concerned by this notification : Yes

Tasks performed by the Body :

Created : 23/03/2007 **| Last update :** 06/05/2008

Product family, product /Intended use/Product range	Procedure/Modules	Annexes or articles of the directives	Limitations
Active medical devices in Class I, IIa or IIb	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	
Non-active medical devices, except Class III 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	