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

From : REPRESENTATION PERMANENTE DE LA FRANCE AUPRES DE L'UNION EUROPEENNE 1000 BRUXELLES

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

Other Member States

Reference : Legislation : 98/79/EC In vitro diagnostic medical devices

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

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Body : **NB 0459**

Created : Unknown (Notifications pre-dating 2006 are not available in these lists) **| Last update :** 27/07/2010

Period of validity of the notification :

Valid until : Unlimited

The body is formally accredited against :

EN 45001

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
In vitro diagnostic medical devices	EC declaration of conformity Full quality assurance system EC type-examination EC verification Production quality assurance	Annex III Annex IV Annex V Annex VI Annex VII	