



Product Service

EC Design Examination Certificate

(Annex IV, section 4 of the Directive 98/79/EC on
In Vitro Diagnostic Medical Devices)

No. V7 06 03 49408 010

Manufacturer: **BAG -
Biologische Analysensystem GmbH**
Amtsgerichtsstr. 1-5
35423 Lich
GERMANY

Product: **Reagents and Reagents products for blood
typing**

Model(s): **Anti-D Blend monoclonal (IgM+IgG)**

Parameters:

Clone:	Id-No.:
-	-
D175-2/D415 1E4	6785 / 6786 / 6787

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the aforementioned devices according to Annex IV, section 4 of the Directive 98/79/EC on In Vitro Diagnostic Medical Devices. The design of the devices conforms to the provisions of this Directive. See also notes overleaf.

Report No.: 70120441

Valid until: 2009-03-31



Date, 2006-04-03

Reiner Krumme

TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 98/79/EC concerning In Vitro Diagnostic Medical Devices with identification no. 0123.

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