



Product Service

EC Design Examination Certificate

(Annex III, section 6 of the Directive 98/79/EC on
In Vitro Diagnostic Medical Devices)

No. V9 07 02 55233 014

Manufacturer: Ulti Med Products (Deutschland) GmbH

Reeshoop 1
22926 Ahrensburg
GERMANY

Product: Self test for detection of blood in stool

Model(s): FOB - Immunological rapid test for
the qualitative detection of human
hemoglobin in stool-self test

Parameters: Modell No.: 010H100

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 III, section 6 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.: 71303244

Valid until: 2012-02-21

Date, 2007-02-21

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.