



Product Service

# EC - CERTIFICATE

## Product Quality Assurance System

(Annex VI of the Directive 93/42/EEC on Medical Devices)

No. G3 08 03 11427 008

**Manufacturer:** LMB Technologie GmbH

Möslstr. 17  
85445 Schwaig  
GERMANY

**Facility(ies):**

LMB Technologie GmbH  
Möslstr. 17, 85445 Schwaig, GERMANY

**Product  
Category(ies):**

**Devices for Treatment of  
Donor Blood and Accessories**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for final inspection and test of the respective products / product categories according to Annex VI, section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class IIb products an additional Annex III certificate is mandatory. See also notes overleaf.

**Report no.:** 71334672

**Valid until:** 2013-05-01



**Date,** 2008-05-01

Reiner Krumme

TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.