



Product Service

EC Certificate

Product Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex VI
(Devices in class IIa or IIb)

No. G3 18 04 11427 020

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 and Cooling
of Donor Blood**

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 devices / device categories in accordance with MDD Annex VI. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report no.:

713129586

Valid from:

2018-05-07

Valid until:

2023-05-01

Date, 2018-05-07

Stefan Preiß



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

Page 1 of 1