



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 08 05 11427 009

Manufacturer: LMB Technologie GmbH

Möslstr. 17
85445 Schwaig
GERMANY

Facility(ies):

LMB Technologie GmbH
Möslstr. 17, 85445 Schwaig, GERMANY

Nanjing Cell-Gene Biomedical Co., Ltd.
No.8 Tianpu Road, Pukou Economic Development Zone, 211800
Nanjing, PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): Blood Bags, Leukocyte Filters,
Infusion Systems, Transfusion Systems

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective products / product categories according to Annex II 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 III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

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Hans-Heiner Junker

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

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