



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 047402 0082 Rev. 00

Manufacturer:

Fresenius Kabi AG

61346 Bad Homburg
GERMANY

Product Category(ies):

ACTIVE MEDICAL DEVICES (class IIa / IIb)

**Devices for Autotransfusion, Apheresis,
Cell Separation, and Photopheresis;**

Accessories for Blood Donation Systems

NON ACTIVE MEDICAL DEVICES (class IIa / IIb)

Transfusion and Transfer Sets incl. Accessories;

Medical Containers; Filters;

Blood Donation Systems and Accessories;

**Disposables for Autotransfusion, Apheresis,
Cell Separation, and Photopheresis**

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 devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

713168590

Valid from:

2020-05-19

Valid until:

2024-05-26

Date,

2020-05-19

Christoph Dicks

Head of Certification/Notified Body