



Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V

(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 17 04 47402 051

Manufacturer:**Fresenius Kabi AG**

61346 Bad Homburg

GERMANY

**Product****Category(ies):****NON ACTIVE MEDICAL DEVICES (class I sterile)****Enteral and Parenteral Feeding Products;****Infusion, Transfusion and Transfer Sets****incl. Accessories; Drainage Products;****Disposables and Accessories for Autotransfusion**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.:

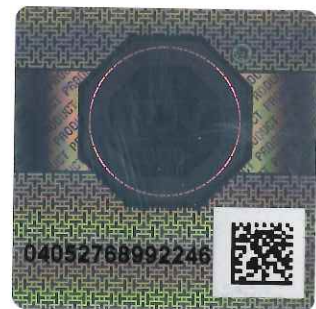
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Valid from:

2017-06-08

Valid until:

2022-06-07



Date, 2017-05-19

Stefan Preiß

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

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Facility(ies):

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Werk Friedberg

Freseniusstrasse 1, 61169 Friedberg,
GERMANY

Fresenius HemoCare Netherlands B.V.

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THE NETHERLANDS

Fresenius Kabi Deutschland GmbH

Werk Bad Hersfeld

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GERMANY

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