



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 024492 2490 Rev. 00**

**Manufacturer:**

**Fresenius Medical Care AG & Co. KGaA**  
61346 Bad Homburg  
GERMANY

**Product Category(ies):**

- Dialysers and Filters for haemodialysis
- Adsorber for therapeutic apheresis
- Tubing systems and accessories for haemodialysis, peritoneal dialysis and apheresis therapies
- Catheters and Accessories for haemodialysis and peritoneal dialysis
- Fistula needles
- Syringes
- Solutions
- Cleaning and disinfectant agents
- Concentrates and solutions for haemodialysis
- Dialysis fluid supply equipment
- Active medical devices for extracorporeal blood treatment and peritoneal dialysis

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.:**

713163830\_1

**Valid from:**

2020-01-22

**Valid until:**

2024-05-26

**Date,**

2020-01-22

Christoph Dicks  
Head of Certification/Notified Body



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**Facility(ies):**

**Fresenius Medical Care AG & Co. KGaA**  
Else-Kröner-Str. 1, 61352 Bad Homburg,  
GERMANY