



Product Service

# EC Certificate

## Production Quality Assurance System

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

No. G2 18 04 14788 026

**Manufacturer:** **Multimedical s.r.l.**  
**Zona Ind. Gerbolina**

Via G. Rossa 69,71,73  
46019 Viadana (MN)  
ITALY



**Facility(ies):** Multimedical s.r.l. Zona Ind. Gerbolina  
Via G. Rossa 69,71,73, 46019 Viadana (MN), ITALY

**Product Category(ies):** Transfusion and infusion sets and associated components: stopcocks, burettes; tubings, extension lines, needles for hemodialysis, paracentesis and thoracentesis, needles for infusion, kit for paracentesis and thoracentesis (needles, syringes and drainage bags), arthroscopy sets, tubing and surgical cannulae for surgical aspiration

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

**Report No.:** ITA1033875

**Valid from:** 2018-05-12

**Valid until:** 2023-05-11



**Date,** 2018-04-17

Stefan Preiß

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

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