



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 14 08 14788 022

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): **Elastomeric infusion pumps**

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.

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Valid until: 2019-09-03



Date, 2014-09-05

Hans-Heiner Junker

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

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