



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 13 06 14788 020

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): Gravity infusion sets and associated components: valves, connectors, adapters, filters, stopcocks, spikes, caps, manifolds, burette; tubings, extension lines, drainage bags, nutrition bags, urine bags, urology sets

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.: ITA 232661

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Hans-Heiner Junker



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

Page 1 of 1