



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 10 31848 018

Manufacturer: **ARIES s.r.l.**
Via XXV Luglio, 43
41037 Mirandola (MO)
ITALY



Facility(ies): ARIES s.r.l.
Via XXV Luglio, 43, 41037 Mirandola (MO), ITALY

Product Category(ies): **Infusion sets for gravity, extension lines, stopcocks, manifolds, accessories for infusion, spikes, valves, infusion sets suitable for angiography, solution filters, nutrition bags, infusion sets with venous pressure manometer, washing sets for urology**

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.: ITA973618

Valid from: 2018-01-14
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Stefan Preiß

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

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