ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 15 03 31848 015

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 with needle, infusion set suitable for photo-dynamic therapy, extension lines, transfusion sets, drainage systems and needles for drainages, infusion sets for pump

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.

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Valid from: 2015-03-16
Valid until: 2018-01-13

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Date, 2015-03-17

Hans-Heiner Junker

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

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