



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 04 38303 026

Manufacturer: Pentaferte Italia S.r.l.

Viale Piane Nocella, 23
64012 Campli (TE)
ITALY



Facility(ies):

Pentaferte Italia S.r.l.
Via Modena 119, 44122 Ferrara, ITALY

Pentaferte Italia S.r.l.
Viale Piane Nocella, 23, 64012 Campli (TE), ITALY

Product

Category(ies):

**Syringes without needle, infusion
sets without needle
and accessories for enteral feeding**

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.: ITA920766

Valid from: 2017-06-02

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Date, 2017-05-31

Stefan Preiß

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

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