



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 026056 0025 Rev. 02

Manufacturer:

Delta Med S.p.a.

Via Guido Rossa 20
46019 Viadana (MN)
ITALY

**Product
Category(ies):**

**Kit of arterial and venous peripheral catheters and
peripheral venous and
arterial catheters with
blood stop system**

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:G2 026056 0025 Rev. 02](http://www.tuvsud.com/ps-cert?q=cert:G2_026056_0025_Rev.02)

Report No.:

ITA 1585257

Valid from:

2021-04-20

Valid until:

2024-05-26

Date,

2021-04-20

Christoph Dicks
Head of Certification/Notified Body