



EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

Manufacturer:

Medesy srl

Viale dell'Industria, 1 - Area industriale

33085 MANIAGO (PN) - ITALIA

Facility(ies)

Medesy srl

Viale dell'Industria, 1 - Area industrial, 33085 MANIAGO (PN) – ITALIA

Product Category(ies):

Dental instruments, laboratory systems and accesories

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 71380125

Valid until: 2022-01-15

Date, 2017-01-21

Hans-Heiner Junker

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

Page 1 of 2

