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Product Service

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)

No. G1 096204 0004 Rev. 01

Manufacturer:

Mesa di Sala Giacomo & C. S.n.c.
Via dell'Artigianato, 35/37
25039 Travagliato (BS)
ITALY

Product Category(ies):

Cobalt, iron, nickel and titanium based alloys for prostheses plates, crowns and bridges, ceramics and soldering; prosthetic components for dental implantology.

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.: ITA10721621SC

Valid from: 2020-03-23
Valid until: 2023-01-16

Date, 2020-03-23

Christoph Dicks
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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Facility(ies): Mesa di Sala Giacomo & C. S.n.c.

Via dell'Artigianato, 35/37, 25039 Travagliato (BS), ITALY

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

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