



Product Service

EC Certificate

Production Quality Assurance

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

No. G2 10 10 40503 011

Manufacturer: 8853 S.p.A.

Via Pitagora, 11
20016 Pero (MI)
ITALY

Facility(ies):

8853 S.p.A.
Via Pitagora, 11, 20016 Pero (MI), ITALY

**Product
Category(ies):**

**Non precious dental alloys,
pigments, porcelains
and resins for use
in dentistry**

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 until: 2016-01-30

Date, 2011-01-31

Hans-Heiner Junker



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

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