



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 16 12 95419 005**

**Manufacturer:** **Quanta System S.P.A.**

Via Acquedotto, 109  
21017 Samarate (VA)  
ITALY

**Facility(ies):**

Quanta System S.P.A.  
Via Acquedotto, 109, 21017 Samarate (VA), ITALY

**Product  
Category(ies):**

**Surgical and therapeutic laser devices,  
therapeutic intense pulsed light devices,  
therapeutic radiofrequency stimulation  
devices, therapeutic devices for  
stimulation of connective tissue,  
combined therapeutic laser and intense  
pulsed light devices, handpieces for  
therapeutic laser and intense pulsed  
light devices, sterile optical fibers for  
surgical laser devices**



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.:** ITA2513872S

**Valid from:** 2017-02-21

**Valid until:** 2020-03-09

**Date,** 2017-02-21

Stefan Preiß



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

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