



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 15 01 90546 002

**Manufacturer:** **Quanta System S.p.A.**  
Via IV Novembre, 116  
21058 Solbiate Olona (VA)  
ITALY

**Facility(ies):** Quanta System S.p.A.  
Via IV Novembre, 116, 21058 Solbiate Olona (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.:** ITA251387

**Valid from:** 2015-03-10

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

Hans-Heiner Junker

**Date,** 2015-02-20



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

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