



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 14 06 11426 016

Manufacturer: LISA laser products OHG

Max-Planck-Str. 1
37191 Katlenburg-Lindau
GERMANY

Facility(ies):

LISA laser products OHG
Max-Planck-Str. 1, 37191 Katlenburg-Lindau, GERMANY

**Product
Category(ies):**

**Therapeutical solid state laser devices for
medical applications and related accessories;
Endoscopes and related instruments, adapters
and consumables**

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.: 713040316

Valid from: 2014-07-23

Valid until: 2019-07-20



Date, 2014-07-24

Hans-Heiner Junker

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

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