



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 019452 0018 Rev. 02

Manufacturer: **Dufner Instrumente GmbH,**

Medizintechnik

Föhrenstrasse 9 78532 Tuttlingen **GERMANY**

Product Category(ies): Endoscopes, Endoscope Shafts,

High Frequency Instruments, Suction Tubes, Suction Valves,

Trocar Sleeves, Trocars,

Insufflation Cannulas, Self-retaining

Retractors and Spreaders

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.: 713169628

Valid from: 2020-03-10

Valid until: 2024-01-13

Date, 2020-03-10

Christoph Dicks

Head of Certification/Notified Body

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

