By Royal Charter

EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

No. **CE 711664**

Issued To: **Geuder AG**
Hertzstraße 4
69126 Heidelberg
Germany

In respect of:

Design, development and manufacture of active, non sterile ocular photocoagulation devices/systems and accessories.

Auslegung, Entwicklung und Produktion von aktiven, nicht sterilen okularen Photocoagulations- Geräten/Systemen und Zubehör.

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):

Gary E Slack, Senior Vice President Medical Devices

First Issued: **2019-08-20** Date: **2019-08-20** Expiry Date: **2024-05-26**

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Supplementary Information to CE 711664

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NBOG Code	Device Name	Intended Purpose per IFU
Class IIb		
MD 1402 MDS 7010	Endotron 532nm surgical system (incl. G-61101 foot switch)	Laser system indicated for use in photocoagulation of both anterior and posterior segments.

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