ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 01 34736 021

Manufacturer: **Candela Corporation**
530 Boston Post Road
Wayland MA 01778
USA

EC-Representative: **Scanlan Group B.V.**
Alasmeerderweg 610
1437 EJ Schiphol-Rozenburg
THE NETHERLANDS

Product Category(ies): **Dermatology Laser System**

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.: DM1201372

Valid from: 2014-03-02
Valid until: 2019-03-01

Date, 2014-02-14

Hans-Heiner Junker

600264

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

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstrasse 65 · 80339 München · Germany

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