



EC-Certificate

SQS as a conformity assessment body identification number 1250 herewith certifies the company

Meridian AG
CH-3608 Thun

the use of a quality assurance system in its design, development, manufacturing and distribution which fulfills the requirements set out in:

ANNEX II.3

Directive 93/42/EEC

This approval is based on the result of the audit documented in the report dated November 20, 2009

The scope of validity covers the products

Ophthalmic Lasers; Microruptor V, Merilas 810Tpro, Merilas 532alpha, SL Laser -532, SL Laser -Nd:YAG, Laserprobe Typ Endoprobe

The following CE label can be applied to these products

CE 1250

A condition for the validity of this certificate is a regular examination in accordance with Annex II.5 of the Directive 93/42/EEC.

Swiss Association for Quality and Management Systems SQS
Bernstrasse 103, CH-3052 Zollikofen
Issue date: November 20, 2009

This SQS Certificate is valid up to and including September 13, 2012
Registration number 32730
Approved Medical Responsible:
November 20, 2009

T. Zahner, Managing Director SQS

K. Schulze, Medical Responsible



SCESm 001
CE 1250



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