



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 10 05 36365 009

Manufacturer: Iskra Medical d.o.o.
Stegne 23
1000 Ljubljana
REPUBLIC OF SLOVENIA

Facility(ies): Iskra Medical d.o.o.
Stegne 23, 1000 Ljubljana, REPUBLIC OF SLOVENIA

Product Category(ies): Medical laser, electro stimulators,
ultrasound equipment, medical products
for lymphdrainage and vacuum therapy,
magnet therapy and phototherapy

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 products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

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Valid until: 2015-05-30

Date, 2010-05-31

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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