



Product Service

# EC - CERTIFICATE

## Production Quality Assurance System

(Annex V of the Directive 93/42/EEC on Medical Devices)

No. G2 09 05 64891 004

**Manufacturer:** LUKROSS INTERNATIONAL S.R.L.  
Via Laboratori Autobianchi, 1 PTB 14/N  
20033 Desio (MI)  
ITALY

**Facility(ies):** LUKROSS INTERNATIONAL S.R.L.  
Via Laboratori Autobianchi, 1 PTB 14/N, 20033 Desio (MI), ITALY

**Product Category(ies):** Electromagnetic stimulation devices

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective products / product categories according to Annex V, 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 IIb and III products an additional Annex III - certificate is mandatory. See also notes overleaf.

**Report No.:** ITA 193926

**Valid until:** 2014-05-31



**Date,** 2009-06-01

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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