

Notified Body No 1023 INSTITUTE FOR TESTING AND CERTIFICATION, Inc.

Zlin, Czech Republic - www.itczlin.cz

EC CERTIFICATE

No. 09 0446 QS/NB

issued in compliance with the Council Directive 93/42/EEC as amended, which is implemented by the Czech Government Order No. 336/2004 (Collection of Laws), certifies that the medical device of Class lib:

Nd: Vaginal tighening laser system

Models: SW-333E-1

manufactured by company

Guangzhou Sunwin Technology Co., Limited No.1 Building, Qishe Intrustrial Park, Tangan Road, Baiyun District, Guangzhou City, China

is manufactured under conditions fulfilling the quality system requirements of Annex II, Section 3.2 of the Directive 93/42/EEC, as amended.

The Notified Body No. 1023 has performed an audit of the above products quality system covering the design, manufacture and final inspection of the certified products. The quality system has been assessed, approved and is subject to continuous surveillance according to Annex II, Sections 3.3 and 5 of the Directive 93/42/EEC. The detailed description of the system parts, requirements and measures applied by the manufacturer are presented in the Final Report No. 803600585A/2009, which is enclosed to this Certificate.

This Gertificate is issued under the following conditions:

- It applies only to the quality system maintained in the manufacture of above referenced models of medical devices and it does not substitute the design or type-examination procedures; if requested.
- The Certificate remains valid until the manufacturing conditions or the quality system are changed but until the 23rd July 2014 at the latest.
- 3. The Certificate validity is conditioned by positive results of surveillance audits.
- 4. After fulfilling the relevant EU legislation, the manufacturer shall affix to each medical device, of the above referenced models, the CE-marking followed by the number of the Notified Body according to this example:

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Issued in Zlin, on 24th July 2012

RNDr. Radomir Čevelik

Representative of the Notified Body No. 1023