

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

Zlin, Czech Republic - www.itczlin.cz

# EC CERTIFICATE

No. 10 0655 QS/NB

issued in compliance with the Council Directive 93/42/EEC as unended, which higherman by the Czeck Government Order No. 336/2004 (Collection of Laws), certified that the medical devices of Classical

### **Derma Roller System**

models:D001,D002,D003,D004,D005,D006,D008,D009

manufactured by company

## Beijing Zhongxing Chuanglian Tech Co. Ltd

No.506, Building 97, Xisanqi, Madian District, Berning, China

are manufactured under conditions. Iffilling the quality system requirements of Annex II, Section 3.2 of the Directive 93/42/5.2C.

The Notified Budy No. 104 has performed a auditor the above products quality system. The quality system has been as essed, all over an as subject continuous supplial and a coording to Annex II, Sections 3.3 and 5 of the Directive 33/42/EE and detailed rescription of the system parts, requirements and measures applied by the magain sture have presented in the Final Report No. 803600930A/2010, which is enclosed to this Certificate.

#### This C rtificate is sever under the following conditions:

- I applies only to be a vality sy im maintained in the manufacture of above referenced models of medical levices and it does not ubstitute the resign or type-examination procedures, if requested.
- The Certificate remains valid use the manufacturing conditions or the quality system are changed but until the 2<sup>nd</sup> August, 2015 pane latest.
- 3. See Certificate validities conditioned by positive results of surveillance audits.
- 4. After 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:

CE1023

1023

RNDr. Radomír Čevelík Representative of the Notified Body No. 1023

Issued in Zlin, on 3rd August 2010