

06/2008



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

Zlin, Chin Republic - www.itczlin.cn

EC CERTIFICATE

No. 09 0169 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 product – medical device of Class Ilb, model

PORTABLE ULTRASOUD 3MHZ

manufacturer

CR s.r.o. vilova 2659, 413 01 Roudnice nad Labem, China Republic

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 product 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. 803600496/2009, which is enclosed to this Certificate.

This Certificate is issued under the following conditions:

- It applies only to the quality system maintained in the manufacture of above referenced model of medical device 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 25th March 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 model, the CE-marking followed by the number of the Notified Body according to this example:

(E₁₀₂₃

Issued in Zlin, on 26th March 2009

RNDr. Radomir Čevelik
Representative of the Notified Body No. 1023