Product Certification Body
INSTITUTE FOR TESTING AND CERTIFICATION, Inc.
Zlin, Czech Republic – www.itczlin.cz

CERTIFICATE

No. 12 0766 T/ITC

Confirms that the product – medical device of Class I according to the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices, as amended, which is implemented by the Czech Government Order No. 336/2004 (Collection of Laws)

Laryngoscopes
Magill Forceps

manufactured by the company

Vision Medical Devices (India) Pvt. Ltd.
Plot no. 586, Industrial area, Phase – 9, Mohali – 160059 (Punjab), INDIA

complies to the applicable essential requirements of the Directive 93/42/EEC.

The ITC Products Certification Body has conducted with successful results the type-examination of the certified product according to the relevant parts of the above mentioned Directive and appropriate harmonized European standards.

The detailed product description, documents, assessment procedures and evaluations of the examination are presented in the Final Report No. 313600324/2012.

Condition of this Certificate use and related information:

1. *It applies only to the above referenced models of the medical devices.*
2. *It does not imply that the ITC has performed any surveillance or control of its manufacture.*
3. *The manufacturer is obligated to assure that all medical devices of the respective models conform to the type approved by this Certificate.*
4. *The Certificate remains valid until the manufacturing conditions, the quality system or relevant legislation are changed but until the **12th September 2015** at the latest.*
5. *After fulfilling the relevant EU legislation requirements, the manufacturer shall affix to each medical device, of the above referenced models, the CE-marking according to this example:*

CE

Issued in Zlín, on 13th September 2012

Institut pro testování a certifikaci, a.s. Zlín

RNDr. Radomír Čevelík
General Director