

144571-15-07-28

EC CERTIFICATE

Full Quality Assurance System

Directive 93/42/EEC on Medical devices, Annex II excluding (4)

CE Certiso Ltd., Organisation for Certification and Testing on the Field of Medical and Hospital Engineering (NB 2409) certifies that the following manufacturer's quality management system concerning to the listed devices and device categories meets the requirements of the related requirements of the directive.

Name of the manufacturer:

Thor Laboratories Kft.

Headquarters: 8000 Székesfehérvár, Raktár utca 2., Hungary

Manufacturing plant: 1119 Budapest, Pajkos utca 50., Hungary

Scope:

Respiratory measurement and diagnostic devices, including spirometers and related software

The certificate covers the following devices:

| Description of the device | Туре | Intended use | Risk class |
|--|--------------|--|---------------|
| PC Spirometer | SpiroTube PC | pulmonary function diagnostics and monitoring | Ila |
| Mobile Edition BlueTooth Spirometer | SpiroTube ME | | IIa |
| Mobile Handheld Spirometer | SpiroThor | | IIa |
| Mobile Handheld Spirometer | OTTHON | | IIa |

This certificate is valid only in case of successfully conducted annual surveillance audits. ID number of the related audit report: 117-CE-150727

Issue: 1

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Start date of certified status: 08 April 2015

Expires:

07 April 2018



CE Certiso
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NB ID number: 2409