

**Full Quality Assurance System**  
Directive 93/42/EEC on Medical devices, Annex II excluding (4)

CE Certiso Ltd. (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:

**Med TeCo LTD.**

Headquarters: **Russia, 141009 Mytishchi, Moscow region, Olympiysky prospect 16\2**  
Authorised representative: **Medical Devices Ltd.  
1149 Budapest Vezér str. 79/2 Hungary**

Scope:

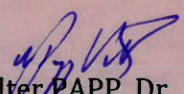
**Universal Plazma Sterilizers**

The certificate covers the following devices:

Description of the device	Intended use	Model	Risk class
Universal Plasma Sterilizer	low temperature hydrogen peroxide plasma sterilization	Plaster-50-Med TeCo	II.b
Universal Plasma Sterilizer	low temperature hydrogen peroxide plasma sterilization	Plaster-100-Med TeCo	II.b
Universal Plasma Sterilizer	low temperature hydrogen peroxide plasma sterilization	Plaster-30C-Med TeCo	II.b

This certificate is valid only in case of successfully conducted annual surveillance audits.  
ID number of the related audit report: 158-CE-170320

Issue: 2  
Issued: 11 September 2019  
First issued: 06 September 2018  
Start date of certified status: 06 September 2018

  
Valter PAPP, Dr.  
General Manager

Expires:  
**05 September 2023**

