

# CERTIFICATE

## for the

### Quality Assurance System



As a notified body of the European Union (Reg. No. 0124) DEKRA Intertek Certification GmbH hereby approves the Quality Assurance System applied for design, manufacture and final inspection by the company

**TELMED Medizintechnik GmbH**  
Breslauer Straße 2 • D – 85386 Eching

Approval is based on the result of the certification audit with report number 50941-Z1-00 and is performed in accordance with the stipulations of

### Annex II, Section 3 of the Directive 93/42/EEC

of the Council dated June 14, 1993 governing medical devices. The certification is applicable to the devices specified in the Annex. The devices in question are subjected to testing and examination in accordance with Annex II, Section 3 of the Directive 93/42/EEC. The listed devices may be affixed with the CE marking indicated below.



Date of the first certification: 11.10.2004

This certificate is valid until: 17.02.2009

Date of the last recertification: ---

Certificate-registration No.: 50941-16-00  
English version

*Thiel*

DEKRA Intertek Certification GmbH  
Stuttgart, 11.10.2004



Akkreditiert durch  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln  
und Medizinprodukten  
ZLG-ZQ-992.94.16

**Annex to the Certificate 50941-16-00 dated 11.10.2004**

English version

Revision status: 1

Date: 03.08.2005

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Devices/device categories included in the certificate

Class II a:

Portable ECG unit, product designation: card-iac-pen M3

Veins monitor, product designation: SVD Monitor COMPACT

