

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:

**MEDITECH Egészségügyi Szolgáltató,
Műszerfejlesztő és Kereskedelmi KFT.**

Headquarters: **1184 Budapest, Mikszáth Kálmán utca 24., Hungary**

Scope:

Ambulatory blood pressure monitors, ECG Holter monitors, combined devices and cardiovascular data management software

This certificate is valid only with the annexes, in case of successfully conducted annual surveillance audits.

ID number of the related audit report: 183-CE-190226

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