

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:

Andramed GmbH.

Headquarters:

Schiesswiesenstr. 18., 72766 Reutlingen

Scope:

Unmounted bare aortic stent

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

ID number of the related audit report: **157-CE-170314**

Issue: 1

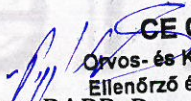
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Start date of certified status: 8 December 2020

Expires:

25 May 2024


CE Certiso
Orvos- és Kórháztechnikai
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