TÜV PROFI CERT

CERTIFICATE

for

DIN EN ISO 13485:2012

(quality managementsystem medical devices)

By means of periodical audits it is assured, that the inspection process assures the product/service conformity including all requirements therefor. This certificate does not acquit the company of its responsibility for the compliance with all legal requirements and service properties.

medifa
Made in Germany

medifa-hesse GmbH & Co. KG
Industriestr. 5
57413 Finnentrop
Germany

scope:

Development, manufacture, distribution and service
of medical-technical devices and furniture

Certificate registration No. **70 105 523**	Certificate valid from 2016-02-15 to **2019-02-14**
Audit report No. 4300 8730	First certification 1998-09-17

Darmstadt, 2016-01-31
Certification body of TÜV Hessen
– Head of Certification body –