

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

scope:

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

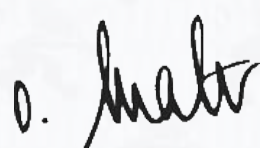
Certificate registration No. **70 105 523**

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 –