

Certificate

Quality Assurance

ecm Zertifizierungsgesellschaft für Medizinprodukte in Europa mbH, Bismarckstr. 106, 52066 Aachen, Germany, hereby declares that an examination of the under mentioned quality assurance system has been carried out following the requirements of DIN EN ISO 13485:2012.



Through an audit performed on behalf of

SMT Schilling Metalltechnik GmbH

Griesweg 33, 78570 Mühlheim, Germany

it could be demonstrated that a quality assurance system

according to **DIN EN ISO 13485:2012**

"Medical devices – Quality management systems – Requirements for regulatory purposes"

for the **development, production and distribution of chirurgical instruments and implants**

has been established and implemented.

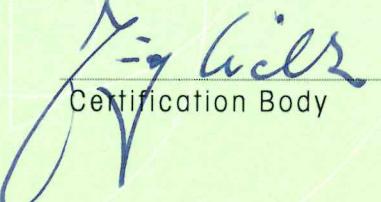
This certificate is only valid under the conditions stated in the hereafter mentioned audit report. Any substantial changes of the quality assurance have to be notified to ECM and are subject to a separate assessment.

Report Number
661-15-422

Registered under
Z/15/03583E

Valid until
June 10th, 2018

Aachen, June 11th, 2015

A handwritten signature in blue ink, appearing to read 'F. G. Lüder'.
Certification Body