

# Certificate

## Quality Management System

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

**IOLUTION GmbH**

Ruhrstraße 13, 22761 Hamburg, Germany

it could be demonstrated that a quality management system

according to

**DIN EN ISO 13485:2012**

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

for the

**development, manufacture and distribution of ophthalmic medical devices / medical devices**

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

217-17-118

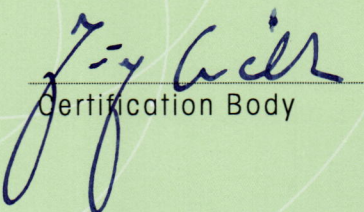
Registered under

Z/17/04133E

Valid until

March 31st, 2019

Aachen, November 2<sup>nd</sup>, 2017

  
Certification Body