



Product Service

CERTIFICATE

No. Q1N 17 11 02403 001

Holder of Certificate: SIFI S.p.A.Via E. Patti 36
95025 Lavinaio - Aci S. Antonio (CT)
ITALY**Facility(ies):**SIFI S.p.A.
Via E. Patti 36, 95025 Lavinaio - Aci S. Antonio (CT), ITALYSIFI S.p.A.
Via E. Patti 34/b, 95025 Lavinaio - Aci S. Antonio (CT), ITALY**Certification Mark:****Scope of Certificate:**

Design and development, production, sales and distribution of intraocular lenses and viscoelastic solutions for ophthalmic applications. Design and development, production and sale of medical devices in liquid and semisolid form for ophthalmic use. Sales under own name of balanced saline solutions for ophthalmic use. Distribution of medical devices for ophthalmic use. Design, development, production, marketing, sales and servicing of equipment and processing diagnostic data and images for ophthalmology

Applied Standard(s):EN ISO 13485:2012 + AC:2012
Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2003 + Cor. 1:2009)
DIN EN ISO 13485:2012
Upgrade required until 2019-03-31

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

Report No.: ITA974824**Valid from:** 2018-01-22**Valid until:** 2020-09-12**Date,** 2018-01-22

Stefan Preiß



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