



Benannt durch/Designated by
 Zentralstelle der Länder
 für Gesundheitsschutz
 bei Arzneimitteln und
 Medizinprodukten
 www.zlg.de
 ZLG-BS-244.10.08



Product Service

EC Certificate

Full Quality Assurance System
 Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
 (Devices in Class IIa, IIb or III)
 (Class IM)

No. G1 060982 0011 Rev. 03

Manufacturer:

**SHANGHAI LINK INSTRUMENTS
CO.,LTD.**

No.1101 Huyi Road, Shanghai, 201802, China

Product Category(ies): Ophthalmic A Scan, Corneal Topographer,
 Vision Screener, Optical Biometer, Fundus
 Camera, Specular Microscope, Rebound
 Tonometer and Delta Ophthalmic
 Ultrasound Scan,Synoptophore
 Auto Refractometer,Auto Lensmeter.

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned
 manufacturer has implemented a quality assurance system for design, manufacture and final
 inspection of the respective devices / device categories in accordance with MDD Annex II. This quality
 assurance system conforms to the requirements of this Directive and is subject to periodical
 surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See
 also notes overleaf.

Report No.: BJ19967022

Valid from: 2020-01-14
Valid until: 2024-05-26

Date, 2020-01-14

C.D.M.

Christoph Dicks
 Head of Certification/Notified Body



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