

Certificate

Standard **ISO 13485:2016**
Certificate Registration No. : **85 105 001 26002**
Audit Report Number : **147190558**

Certificate Holder:

AUROLAB

No. 1, Sivagangai Main road,
Veerapanjan, Madurai 625020, India

Scope:

Design and Development, Manufacture and Distribution of Sterile Medical Devices Such as:

- PMMA Intraocular Lenses,
- Hydrophobic Foldable Intraocular Lenses,
- Hydrophilic Foldable Intraocular Lenses,
- Hydrophilic Foldable Intraocular Lenses with Injector & Cartridge,
- Preloaded Hydrophobic Foldable Intraocular Lenses,
- Capsular Tension Rings,
- Ophthalmic Solutions,
- Non Absorbable Ophthalmic Suture with Needles,
- Micro Surgical Suture with Needles,
- Absorbable Ophthalmic Suture with Needles,
- Polyglycolic Acid,
- Ptosis Slings,
- Glaucoma Shunt,
- Ophthalmic Surgical Blades,
- Ophthalmic Surgical Instruments.

Note: Refer Addendum.



Bangalore
2026-05-24

The Certification Body of
TÜV Rheinland (India) Pvt. Ltd.
27/B, 2nd Cross Road,
Electronic City, Phase-I,
Bangalore – 560 100, India.
CIN: U72501KA1996PTC020653

MS-0042439_Annexure5

Revision: 0

ADDENDUM

Certificate

Standard **ISO 13485:2016**
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Manufacturing and Distribution of Sterile Medical Devices:

- Injectors Cartridges and Cartridges for injecting Foldable Lenses.

Manufacturing, Distribution, Installation and Service of Active Ophthalmic Medical Equipment such as

- Green laser Photo Coagulator,
- Digital Vision Chart,
- Autorefractor

Proof has been furnished by means of an audit that the requirements of ISO 13485:2016 are met.

The due date for Surveillance Audits is (05-21)

Validity:

The certificate is valid from 2026-05-24 until 2029-05-23.



03/NABCB/00

The validity of this certificate is subject to timely completion of Surveillance audits as agreed in the contract.
The validity of the certificate can be verified under www.tuv.com with the Identification No. 000000000

MS-0042439_Annexure5

Revision: 0