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Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
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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)

**No. G1 095857 0017 Rev. 00**

**Manufacturer:**

**HOYA Medical Singapore Pte. Ltd.**

455A, Jalan Ahmad Ibrahim

Singapore 639939

SINGAPORE

## Product Category(ies): Sterile Preloaded Intraocular Lenses

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:G10958570017Rev.00](http://www.tuvsud.com/ps-cert?q=cert:G10958570017Rev.00)

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2021-04-12

**Valid until:**

2024-05-26

**Date,**

2021-04-12

Christoph Dicks

Head of Certification/Notified Body