



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 103773 0005 Rev. 01

Manufacturer:

CooperVision Manufacturing Limited

South Point

Hamble

Southampton, SO31 4RF

UNITED KINGDOM

Product Category(ies): Non-Active Ophthalmologic Devices

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:G11037730005Rev.01

Report No.:

75947288 - 75950871

Valid from:

2021-05-10

Valid until:

2024-05-26

Date,

2021-05-10

Christoph Dicks

Head of Certification/Notified Body



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Soft (Hydrophilic) Contact Lens Products for the Correction of Ametropia

Soft (Hydrophilic) Contact Lens Products for the Control of Myopia

Soft Contact Lens Care Products

Rigid Gas Permeable Contact Lens Care products

Ancillary Products :

Comfort drops

Preservative free saline