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)

No. G1 020543 0017 Rev. 01

Digitally signed by Bolea Petru
Date: 2021.10.08 12:22:44 EEST
Reason: MoldSign Signature
Location: Moldova

Manufacturer:

Morcher GmbH
Kapuzinerweg 12
70374 Stuttgart
GERMANY

Facility(ies):

Morcher GmbH
Kapuzinerweg 12, 70374 Stuttgart, GERMANY

Product Category(ies):

Intraocular Lenses
Capsular Rings
Pupil Dilator
Injectors for Eye Implants

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.: 713169392

Valid from: 2020-01-24
Valid until: 2024-05-26

Date, 2020-01-24

Christoph Dicks
Head of Certification/Notified Body

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

A4 / 07.17

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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