EU Certificate

Quality Management System
REGULATION (EU) 2017/745 on Medical Devices
Annex IX Chapters I and III

Registration No.:	HZ 1594091-1
Manufacturer:	Carl Martin GmbH Neuenkamper Str. 80-86 42657 Solingen Germany
EUDAMED Single Registration No.:	DE-MF-000005066

Products:

Products of class I, reusable surgical instruments:

L0205 - NEEDLE HOLDERS, REUSABLE
L0901 - ORTHOPAEDIC SURGERY SPOONS AND CURETTES, REUSABLE
L0399 - GENERAL SURGERY INSTRUMENTS, REUSABLE - OTHER
L0904 - OSTEOTOMES AND SCALPELS, REUSABLE
L031399 - GENERAL SURGERY FORCEPS, REUSABLE - OTHER
L031199 - PROBES AND MIRRORS, REUSABLE – OTHER
L010499 - SURGICAL SCISSORS, REUSABLE – OTHER

The Notified Body hereby declares that the requirements of Annex IX, Chapter I of the REGULATION (EU) 2017/745 have been met for the listed products. The above named manufacturer has established and applies a quality management system, which is subject to periodic surveillance, defined by Annex IX, Chapter I, Section 3 of the aforementioned regulation. The requirements of Annex IX, Chapter III are fulfilled.
If class III devices or class IIb implantable devices referred to in the second subparagraph of Article 52(4) are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 4.9 is required before placing them on the market.

Report No.:	1206333-10
Effective date:	2026-06-30
Expiry date:	2031-06-29
Issue date:	2026-06-25

Dipl.-Ing. Anja Fechner
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

This certificate can be validated on https://www.certipedia.com

TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/745 concerning medical devices with the identification number 0197.

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-091

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Neuenkamper Str. 80-86
42657 Solingen
Germany

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Products:

L031308 - SURGICAL AND ANATOMICAL TWEEZERS, REUSABLE
L010101 – MONOBLOCK SURGICAL SCALPELS, REUSABLE

The scope of certification is limited to the aspects relating to the reuse of the device, in particular
cleaning, disinfection, sterilization, maintenance and functional testing and the related instructions for use

Authorized representative(s): N/A

Certificate history		
Revision:	Description:	Issue date:
1	Re-certification Replace certificate HZ 1594091-1 rev. 0 issued 2021-11-04	2026-06-24
2	Re-certification Replace certificate HZ 1594091-1 rev. 1 issued 2021-11-04 Correction of Revision	2026-06-25

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