



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
BS-MDR-099



Product Service

EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III
(Class I Devices in sterile condition, with measuring function or reusable surgical instruments)

No. G11 031564 0016 Rev. 00

Manufacturer:

Ulrich AG

Mövenstraße 12
9015 St. Gallen
SWITZERLAND

The Certification Body of TÜV SÜD Product Service GmbH certifies that the manufacturer has established, documented and implemented a quality management system as described in Article 10 (9) of the Regulation (EU) 2017/745 on medical devices. Details on device categories covered by the quality management system are described on the following page(s).

The Report referenced below summarises the result of the assessment and includes reference to relevant CS, harmonized standards and test reports. The conformity assessment has been carried out according to Annex IX Chapter I and III of this regulation with a positive result. As applicable the involvement of the notified body is limited to the aspects relating to:

- establishing, securing and maintaining sterile conditions,
- conformity of the devices with the metrological requirements,
- reuse of the device, in particular cleaning, disinfection, sterilization, maintenance and functional testing and the related instructions for use.

The certified quality management system is subject to periodical surveillance by TÜV SÜD Product Service GmbH.

Report No.: 713179391

Valid from: 2020-09-01

Valid until: 2025-07-20

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2020-09-01



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No. G11 031564 0016 Rev. 00

Classification: I

Device Group: MDN 1208 - Non-active non-implantable instruments

Device Properties: MDS 1006 - Reusable surgical instruments

The validity of this certificate ./.
depends on conditions and/or
is limited to the following: