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

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 012163 0089 Rev. 02

Manufacturer:

seca gmbh & co. kg
Hammer Steindamm 3-25
22089 Hamburg
GERMANY

SRN Manufacturer - DE-MF-000005469

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.
All applicable requirements of the Testing, Certification, Validation and Verification Regulations TÜV SÜD Group have to be complied with.
For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G11 012163 0089 Rev. 02

Report No.: 713367772

Preceding Certificate No.: G11 012163 0089 Rev. 01

Valid from: 2025-08-12
Valid until: 2030-08-11

Date of Initial Issuance: 2020-08-12

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2025-07-25

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123
TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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

TÜV SÜD 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 012163 0089 Rev. 02

Classification:	Class I
Device Group:	V0399 - MEASUREMENT DEVICES - OTHER
Device Properties:	MDS 1010 - Devices with a measuring function

Classification:	Class I
Device Group:	V030209 - CLINICAL ALTIMETERS
Device Properties:	MDS 1010 - Devices with a measuring function

Classification:	Class I
Device Group:	V030210 - WEIGHING SCALES (EXCLUDING THOSE FOR NEONATOLOGY)
Device Properties:	MDS 1010 - Devices with a measuring function

Classification:	Class I
Device Group:	V030211 - MULTIDIMENSIONAL MEASUREMENTS STATIONS
Device Properties:	MDS 1010 - Devices with a measuring function

Classification:	Class I
Device Group:	Z12080402 - NEONATAL SCALES
Device Properties:	MDS 1010 - Devices with a measuring function

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

Revision History:

Rev.	Dated	Report	Description
00	2020-08-12	713176915	-
01	2021-05-05	713176915	-
02	2025-08-12	713367772	Renewal of certificate

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123
TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

TÜV®