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

Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapter I

Certificate No. G15 019931 0023 Rev. 00

Manufacturer:

GS Elektromedizinische Geräte G. Stemple GmbH
Hauswiesenstr. 26
86916 Kaufering
GERMANY

SRN Manufacturer - DE-MF-000007648

The quality management system has been evaluated in accordance with Regulation (EU) 2017/745, Annex IX Chapter I with a positive result.

Details on devices covered by the quality management system are described on the following page(s). The report referenced below summarises the results of the assessment and includes reference to relevant CS, harmonised standards and test reports.
The certified quality management system is subject to periodical surveillance.

If class I devices in sterile conditions, with measuring function, or reusable surgical instruments are covered by this certificate, the audit was limited to the respective 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.

If class IIa or class IIb devices are covered by this certificate, the quality management system assessment was accompanied by the assessment of technical documentation for devices selected on a representative basis. The periodical surveillance includes further assessment of the technical documentation on the basis of representative samples.
If class III or class IIb implantable devices are covered by this certificate, an EU Technical Documentation Assessment Certificate in accordance with Annex IX Chapter II is required before placing them on the market.

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:G15 019931 0023 Rev. 00

Report No.:	713371610/713371609
Preceding Certificate No.:	G10 019931 0015 Rev. 01 G12 019931 0020 Rev. 00
Valid from:	2025-11-19
Valid until:	2030-11-18

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2025-09-12

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Certificate No. G15 019931 0023 Rev. 00

Classification:	Class III
Device Group:	MDA 0305 - Active non-implantable devices for stimulation or inhibition
Intended Purpose:	See product certificate

Classification:	Class IIb
Device Group:	Z12030282 - VITAL SIGNS MONITORING INSTRUMENTS - SOFTWARE ACCESSORIES
Intended Purpose:	Telemetry application for analyzing medical data and obtaining a teleconsile.

Classification:	Class IIb
Device Group:	Z120304 - CARDIAC COMPRESSORS
Intended Purpose:	The corpuls cpr is a device for electro-mechanical chest compressions within the framework of a cardio-pulmonary resuscitation.

The validity of this certificate depends on conditions and/or is limited to the following:	-none-

Revision History:

Rev.	Dated	Report	Description
00	2025-11-19	713371610/713371609	Renewal of certificate

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