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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 075302 0063 Rev. 00

Manufacturer:

Devicor Medical Products, Inc.

Fifth Floor
300 E-Business Way
Cincinnati OH 45241
USA

SRN Manufacturer:

US-MF-000003322

**Authorized
Representative:**

CEpartner4U BV
Esdoornlaan 13, 3951 DB Maarn, THE NETHERLANDS

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 and certification regulation of TÜV SÜD Group have to be complied with.

For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G11_075302_0063_Rev.00

Report No.:

713203734

Valid from:

2022-09-20

Valid until:

2027-09-19

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2022-09-20



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Classification: I
Device Group: A019012 - BREAST NODULES AND OTHER STRUCTURES
LOCALISATION NEEDLES AND SYSTEMS
Device Properties: MDS 1005.2 - Sterilisation by irradiation

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