



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 045286 0073 Rev. 02

Manufacturer:

**Lohmann & Rauscher
International GmbH & Co. KG**

Westerwaldstraße 4
56579 Rengsdorf
GERMANY

Product Category(ies): Swabs and balls, active medical devices for negative pressure wound therapy including accessories, wound dressings, surface disinfectants, surgical single use instruments

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

713152842/713160691

Valid from:

2020-05-27

Valid until:

2024-05-26

Date,

2020-05-27

Christoph Dicks
Head of Certification/Notified Body

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Surgical Instruments

IIa

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