





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**

Productgroup/ Product name	Classification	Rule
<b>Swabs and Balls</b>	<b>IIa,</b>	<b>6, 7</b>
<b>Active medical devices for negative pressure wound therapy incl. Accessories</b>	<b>IIa, IIb</b>	<b>4, 11, 8</b>
<b>Wound Dressings</b>	<b>IIa, IIb, III</b>	<b>4, 13, 17</b>
Alginate Dressings	IIb, III	4, 13
Hydrocolloid Dressings	IIa, IIb	4
Hydroactive Fiber	IIb	4
Hydrobalance Wounddressings	IIb, III	4, 13
Film Dressings	IIa	4
Contact Net	IIb	4
Foam Dressings	IIb	4
Gel Dressings	IIb	4
Activated Charcoal Dressings	IIb	4
Vaseline Dressings	IIb	4
Superabsorbent Dressings	IIb	4
Aluminized Dressings	IIb	4
Collagen Wound Dressings	III	17
Dressing Material, impregnated with Iodoform	III	13
<b>Surfacedisinfekt</b>	<b>IIa</b>	<b>15</b>

Page 2 of 3

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®

TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD  
ZERTIFIKAT ◆ CERTIFICATE ◆ 認證書 ◆ CERTIFICADO ◆ CERTIFICAT

A4 / 07.17



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**

**Surgical Instruments**

**IIa**

**6**





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

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V

(Devices in class I in sterile conditions, sterilised systems or procedure packs)

**No. G2S 045286 0075 Rev. 01**

## Manufacturer

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

Westerwaldstraße 4  
56579 Rengsdorf  
GERMANY

## Product Category(ies):

**Procedure Packs/ set systems, swabs and balls,  
ophthalmological devices, wound dressings, OR-  
materials, products for compression/ retention and  
support, material for padding, maternity pads,  
accessories to active medical devices for negative  
pressure wound therapy, ENT-devices**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

**Report No.:** 713163955/713157373

**Valid from:** 2020-05-29

**Valid until:** 2024-05-26

**Date,** 2020-05-29

Christoph Dicks  
Head of Certification/Notified Body

TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD  
ZERTIFIKAT ♦ CERTIFICATE ♦ 認證書 ♦ CERTIFICADO ♦ CERTIFICAT



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
[www.zlg.de](http://www.zlg.de)  
ZLG-BS-244.10.08



Product Service

# EC Certificate

Production Quality Assurance System  
Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

**No. G2S 045286 0075 Rev. 01**

Productgroup	Classification	Rule
<b>Procedure Packs/ Set Systems</b>	<b>Art. 12</b>	<b>---</b>
<b>Swabs and Balls</b>	<b>Is</b>	<b>4</b>
<b>Opthalmological Devices</b>	<b>Is</b>	<b>4, 1</b>
<b>Wound Dressings</b>	<b>Is</b>	<b>4, 1</b>
<b>OR-Materials</b>	<b>Is</b>	<b>1, 4</b>
<b>Products for compression, retention and support</b>	<b>Is</b>	<b>1</b>
<b>Material for padding</b>	<b>Is</b>	<b>1</b>
<b>Maternity Pads</b>	<b>Is</b>	<b>1</b>
<b>Accessories to active medical devices for negative pressure wound therapy</b>	<b>Is</b>	<b>2, 4, 1</b>
<b>ENT Devices</b>	<b>Is</b>	<b>5</b>

TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD  
ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ CERTIFIKAT ◆ CERTIFICADO ◆ CERTIFICAT



TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD  
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證書 ◆ CERTIFICADO ◆ CERTIFIKAT ◆ CERTIFICATE



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

Production Quality Assurance System  
 Directive 93/42/EEC on Medical Devices (MDD), Annex V  
 (Devices in class I in sterile conditions, sterilised systems or procedure packs)

**No. G2S 045286 0075 Rev. 01**

## Manufacturer

**Lohmann & Rauscher  
 International GmbH & Co. KG**  
 Westerwaldstraße 4  
 56579 Rengsdorf  
 GERMANY

## Product Category(ies):

**Procedure Packs/ set systems, swabs and balls,  
 ophthalmological devices, wound dressings, OR-  
 materials, products for compression/ retention and  
 support, material for padding, maternity pads,  
 accessories to active medical devices for negative  
 pressure wound therapy, ENT-devices**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

**Report No.:** 713163955/713157373

**Valid from:** 2020-05-29

**Valid until:** 2024-05-26

**Date,** 2020-05-29

Christoph Dicks  
 Head of Certification/Notified Body

