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 057666 0061 Rev. 00

Manufacturer:

Ethicon Endo-Surgery, LLC
475 Calle C
00969 Guaynabo
PUERTO RICO USA

EC-Representative:

Ethicon Endo-Surgery (Europe) GmbH
Hummelsbütteler Steindamm 71, 22851 Norderstedt, GERMANY

Product Category(ies):

**Surgical Devices for Instrument Access, Cutting, Stapling and Suturing;
Ultrasonic Surgical Devices and Accessories;
Electrosurgical Devices and Accessories;
Instruments for Minimally Invasive Endoscopic Procedures
(Surgical or through natural body orifices), except Class I devices
(non-sterile and/or without measuring function)**

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.: 713157818

Valid from: 2019-07-01
Valid until: 2024-05-26

Date, 2019-06-17

Stefan Preiß
Head of Certification/Notified Body

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Facility(ies):

Ethicon Endo-Surgery, LLC
475 Calle C, 00969 Guaynabo, PUERTO RICO USA

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TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany