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

EC Design-Examination Certificate
Directive 93/42/EEC on Medical Devices (MDD), Annex II (4)
(Devices in Class III)

No. G7 057666 0049 Rev. 01

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

Product: **Non-Active Implants**
Surgical Staple, nonbioabsorbable

Model(s): **ENDOPATH ECHELON™ Vascular Reload**

Parameter: ENDOPATH ECHELON™ Vascular Reload:
VASECR35

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the respective devices in accordance with MDD Annex II (4). The design of the devices conforms to the requirements of this Directive. For marketing of these devices an additional Annex II certificate is mandatory. See also notes overleaf.

Report no.: 713150827

Valid from: 2020-02-25
Valid until: 2024-05-26

Date, 2020-02-25

Christoph Dicks
Head of Certification/Notified Body

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

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