



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 0050 Rev. 01**

**Manufacturer:**

**Ethicon Endo-Surgery, LLC**

475 Calle C  
00969 Guaynabo  
PUERTO RICO USA

**Product:**

**Non-Active Implants**

**Model(s):**

**Linear Cutter Reloads**

**Parameter:**

Variants	Model No.
Selectable 55mm	SR55
Selectable 75mm	SR75

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.:** 713150828

**Valid from:** 2020-03-30

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

**Date,** 2020-03-30

Christoph Dicks  
Head of Certification/Notified Body