



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 15 06 57666 050

**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:** **Non-Active Implants**

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.:** 713057450

**Valid from:** 2015-07-13

**Valid until:** 2020-07-12

Hans-Heiner Junker

**Date,** 2015-07-01

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

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**Model(s):** **Linear Cutter Reloads**

<b>Parameters:</b>	<b>Variants</b>	<b>Model No.</b>
	Selectable 55mm	<b>SR55</b>
	Selectable 75mm	<b>SR75</b>

**Facility(ies):** Ethicon Endo-Surgery S.A. de C.V., Avenida de Las Torres 7125  
Colonia Salvarcar 118, 32580 Ciudad Juarez, Chihuahua,  
MEXICO

Ethicon Endo-Surgery, Inc.  
3801 University Boulevard SE, Albuquerque, NM 87106, USA

**Design Facility(ies):** Ethicon Endo-Surgery, Inc.  
4545 Creek Road, Cincinnati OH 45242, USA