

## Declaration of Conformity (DoC)

**Manufacturer:**

**Name:** Ethicon Endo-Surgery, LLC  
**Address:** 475 Calle C  
Guaynabo, Puerto Rico 00969 USA

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**Authorized Representative:**

**Name:** Ethicon Endo-Surgery (Europe) GmbH  
**Address:** Hummelsbuetteler Steindamm 71  
22851 Norderstedt, Germany

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**Products/Model No:**

The following ECHELON ENDOPATH Stapler Reloads and ENDOPATH ECHELON Reloads product codes are CE marked:

**ECR60B, ECR60D, ECR60G, ECR60W, ECR45B, ECR45D, ECR45G, ECR45W, ECR45M, ECR60M, GST60W, GST60B, GST60D, GST60G, GST60T, GST45W, GST45B, GST45D, GST45G, GST45T**

(MDD applicable only)

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**Directive 93/42/EEC declaration (MDD):**

We, Ethicon Endo-Surgery, being the manufacturer, declare that the products listed above meet the applicable provisions of the Medical Device Directive 93/42/EEC including applicable harmonized standards listed in the above referenced Design Dossier.

Products are classified as Class III, Rule 8, per Annex IX.

In order to affix CE marking under this directive, Ethicon Endo-Surgery, has followed the procedure relating to the EC declaration of conformity set out in MDD Annex II excluding (4) and Annex II (4). The full quality system has been certified by TÜV SÜD Product Service GmbH, a Notified Body authorized to carry out such assessments and having the designation 0123 (G1 057666 0061 Rev.00, G7 057666 0054 Rev.01).

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This declaration of conformity is issued under the sole responsibility of the manufacturer.

**Approved by:**

Nathan Anderson  
Sr. Director Quality Make  
Ethicon Endo-Surgery, LLC



(Signature)

21 JUL 2020

(Date)