

# EU Technical Documentation Assessment Certificate

Regulation (EU) 2017/745, Annex IX Chapter II

## MDR 756446 R000

**Manufacturer:** Apollo Endosurgery, Inc.

**Address:**

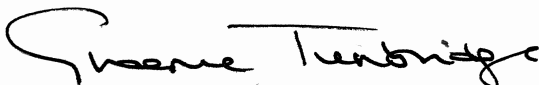
1120 Capital of Texas Highway  
Building 1, Suite 300  
Austin  
Texas  
78746  
USA

**Single Registration Number:** US-MF-000003488

**Scope:** See attached **Device Schedule**

On the basis of our assessment of the technical documentation in accordance with Regulation (EU) 2017/745, Annex IX Chapter II, the technical documentation meets the requirements of the Regulation. For the placing on the market of these devices an additional Annex IX Chapter I and III certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):



Graeme Tunbridge, Senior Vice President Medical Devices

First Issue Date: **2022-11-16**

Current Issue Date: **2022-11-16**

Starting Validity Date: **2022-11-16**

Expiry Date: **2027-11-15**

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### Device Schedule:

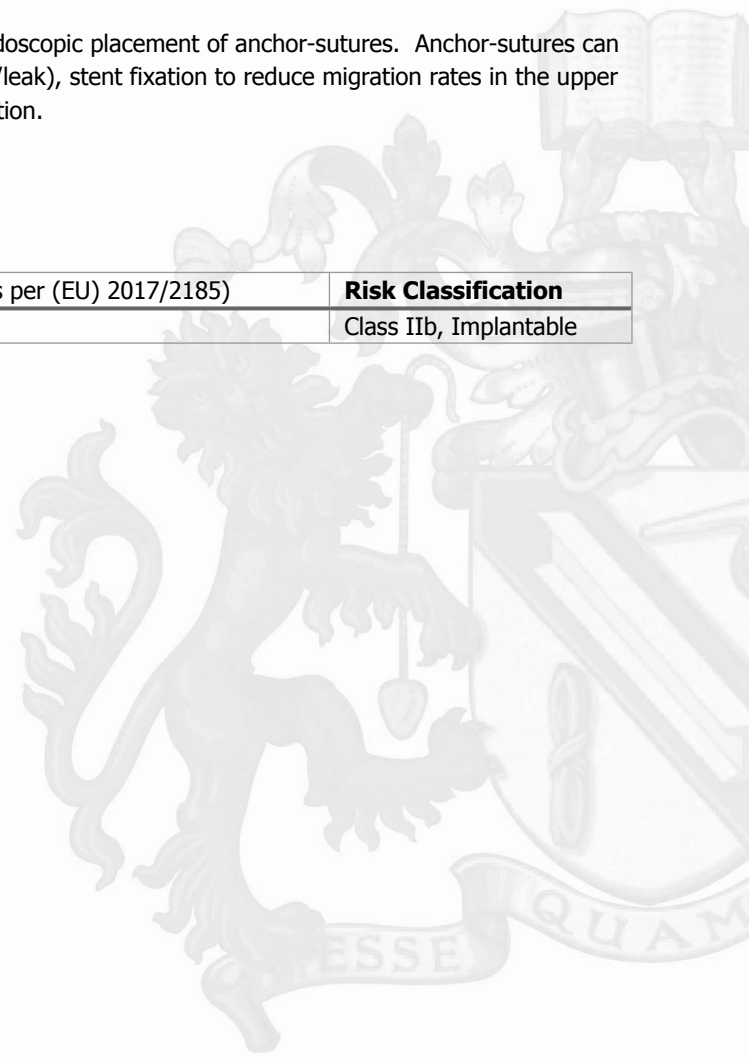
### Intended Purpose as per the Instructions for Use:

The OverStitch™ Endoscopic Suture System is indicated for the endoscopic placement of anchor-sutures. Anchor-sutures can be placed for defect closure (e.g. perforation, ESD/EMR and fistula/leak), stent fixation to reduce migration rates in the upper GI tract, endoscopic sleeve gastropasty, and transoral outlet reduction.

**Risk Classification:** Class IIb, Implantable

**Basic UDI-DI:** 081195502381337C8

Device Name	Model	Type (Codes as per (EU) 2017/2185)	Risk Classification
OverStitch™ Suture Cinch	CNH-G01-000	MDN 1104	Class IIb, Implantable



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### Certificate History

*(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate.Verification@bsigroup.com)*

Date	Reference Number	Action
Current	3507242	Issued



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Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.  
This certificate was issued electronically and is bound by the conditions of the contract.

NB Contact: BSI Group The Netherlands B.V., Say Building, John M. Keynesplein 9, 1066 EP, Amsterdam, Netherlands. Tel: + 31 (0) 20 346 07 80  
Corporate Contact: BSI Group Assurance Limited, registered in England under number 05435540 at 389 Chiswick High Road, London, W4 4AL, UK.  
A Member of the BSI Group of Companies.