EU Quality Management System Certificate

Regulation (EU) 2017/745, Annex IX Chapter I and III

MDR 719610 R000

Manufacturer: Ethicon, LLC

Address:
475 C Street, Los Frailes Industrial Park
Suite 401
Guaynabo
Puerto Rico
00969
USA

Single Registration Number: US-MF-000013111

EU Authorised Representative: Johnson & Johnson Medical GmbH

Address:
Robert-Koch-Strasse 1
Norderstedt
22851
Germany

Scope: See attached **Device Schedule**

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/745, Annex IX Chapter I and III, the quality system meets the requirements of the Regulation. For the placing on the market of Class III devices, and Class IIb implantable devices that are not considered well-established technologies as specified in Article 52(4) an additional Annex IX Chapter II 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: **2021-10-14**

Current Issue Date: **2023-07-05**

Starting Validity Date: **2023-07-05**

Expiry Date: **2026-10-13**

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Device Schedule: Class III and Class IIb devices

Class III, Implantable	**Intended purpose**
ETHICON SECURESTRAP™ Family of Products	See MDR 719665
MERSILENE™ Suture	See MDR 719649
MONOCRYL™ Suture	See MDR 719654
VICRYL RAPIDE™ Suture	See MDR 719664
ETHIBOND EXCEL™ Suture	See MDR 719647
VICRYL™ Suture	See MDR 719663
ETHILON™ Suture	See MDR 719648
PROLENE™ Suture	See MDR 719658
PERMA-HAND™ Braided Silk and Virgin Silk Non-Absorbable Sutures	See MDR 719650
Class IIb, Implantable, Well-established technologies	**Intended purpose**
Stainless Steel Suture	Stainless Steel Suture is indicated for use in sternal closure and orthopedic procedures.

Device Schedule: Class IIa, Custom-made and other devices

Device(s)	**Risk Classification**
Suturing Device	Class Ir

For Class Ir devices (Class I re-usable surgical instruments), the Notified Body conformity assessment is limited to the aspects relating to the reuse of the device.

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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
2021-10-14	3091327	Issued
2023-01-13	3736760	Supplemented – Addition of MERSILENE Suture, MONOCRYL Suture, VICRYL RAPIDE Suture, ETHIBOND EXCEL Suture and VICRYL Suture Amended – Addition of Legal Manufacturer Single Registration Number Amended – Administrative correction to Legal Manufacturer and EU Authorised Representative address format
2023-03-08	3814853	Supplemented – Addition of Coated VICRYL Plus Antibacterial Suture
Current	3908217	Supplemented – Addition of ETHILON Suture, PROLENE Suture, PERMA-HAND Braided Silk and Virgin Silk Non-Absorbable Suture, and Stainless Steel Suture.

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