



EU Quality Assurance Certificate

Regulation (EU) 2017/745, Annex XI Part A

MDR 725202 R000

Manufacturer: 3M Company

Address:

2510 Conway Ave. Saint Paul Minnesota 55144 USA

Single Registration Number: US-MF-000014086

EU Authorised Representative: 3M Deutschland GmbH

Address:

Healthcare Business Carl-Schurz-Str. 1 41443 Neuss Germany

Scope: See attached Device Schedule

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/745, Annex XI part A, the quality system meets the requirements of the Regulation. For the placing on the market of Class III and Class IIb devices an additional Annex X 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 Global Regulatory & Quality

First Issue Date: **2021-02-08** Starting Validity Date: **2025-04-04**

Current Issue Date: **2025-04-04** Expiry Date: **2026-02-07**

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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.





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

Device(s)	Risk Classification	
Sterile disinfecting port protector devices	Class IIa	4
Barrier film	Class IIa	
Warming Blankets	Class Is	
Self-Adherent Wrap	Class Is	7
Skin Closures	Class Is	
Surgical Drapes	Class Is	12
Incise drapes	Class Is	
Surgical drapes – other	Class Is	
Skin staple removers	Class Is	10
Barrier Film	Class Is	
Intravascular protection devices	Class Is	

For Class Is devices, the Notified Body conformity assessment is limited to the aspects relating to establishing, securing and maintaining sterile conditions.

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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
2021-02-08	3154149	First issue.
2021-04-18	3424211	Supplemented – Addition of device group Class Is
2022-03-02	3486382	Supplemented – Addition of device group Barrier film and subcontractor Amended – Addition of Legal Manufacturer SRN number. Amended – Administrative change to correct EU representative address zip code Amended – Administrative change to remove 3M Company site. Amended – Subcontractor change Amended – Subcontractor address updated. Amended – Administrative error, Subcontractor missing after cert reissue. Subcontractor readded. Amended – Administrative, previous certificate history (Ref 3424211) missing amendment for addition of
		subcontractors
2022-07-13	3699536	Amended – Removal of subcontractors
2024-02-20	30107696	Amended – Addition of Ethylene oxide sterilisation subcontractor. Amended – removal of subcontractor names from certificate history.
2024-09-04	30232707	Amended – Removal of subcontractor and addition of new subcontractor.
Current	30380337	Amended – Addition of Ethylene oxide sterilisation subcontractor.

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