

EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Company Single Registration Number: US-MF-000014086 2510 Conway Ave. St. Paul, MN 55144 USA

hereby declare under our sole responsibility that the following CE marked device(s)

Trade Name	3M [™] Transpore [™] Surgical Tape
Intended	Surgical Tape
Purpose	
Reference	1527-0, 1527-1, 1527-2, 1527-3, 1527-08, 1527-18, 1527-
	2S, 1527S-1, 1527S-2, 1527(Bulk), 1527P-2S, 1527NP-
	1S, 1527IP-1SD, 1527P-1SD, 1527NP-1SD, 1527P-1SD
Basic UDI-DI	06082238401010000000015A7

are classified per rule 1 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class I devices in accordance with all applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.

EU Authorized Representative:

EU Representative Address 3M Deutschland GmbH Health Care Business DE-AR-000011642 Carl-Schurz-Str. 1 41453 Neuss, Germany

DocuSigned by: Nadia Battah	8/9/2023
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Nadia Battah	Location/Date
Regulatory Affairs Manager	

3M is a trademark of 3M.

3M Medical Solutions Division