

**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™ Paper Surgical Tape 3M™ Micropore™ Surgical Tape
Intended Purpose	Surgical Tape
Reference	840W-05, 840W-15, 840W-1, 840W-2 1533IP-2S, 1533IP-1S, 1533NP-1S, 1533IP-0S, 1533NP-0S, 1533SP-1, 1533P-1SD, 1533NP-0SD, 1533NP-1SD, 1533SP-1D, 1533SP-0D, 1533P-0D, 1533P-1D
Basic UDI-DI	06082238401010000000058AR

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
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DocuSigned by:

Dianne Gibbs

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11/16/2022

Dianne Gibbs, RAC
Regulatory Affairs Director
3M Medical Solutions Division

Location/Date

3M is a trademark of 3M.