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™ Micropore™ Surgical Tape
Intended Purpose	Surgical Tape
Reference	1533-0, 1533-1, 1533-2, 1533 (Bulk)
Basic UDI-DI	06082238401010000000009S

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:
Dianne Gibbs
8146ED4E5FCD4D1...

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

11/10/2022
Location/Date