

**EUROPEAN MEDICAL DEVICE REGULATION****Declaration of Conformity**

As Legal Manufacturer, we

3M Company
Single Registration Number (TBD)
2510 Conway Ave. St. Paul, MN 55144 USA

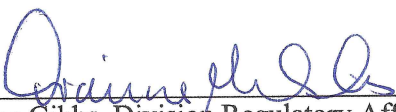
hereby declare under our sole responsibility that the following CE marked device(s)

Trade Name	Transpore™ Surgical Tape
Intended Purpose	A general-purpose tape for the hospital and home care patient used to secure most dressings, tubing and devices to skin.
Catalogue Number	1527-0, 1527-1, 1527-2, 1527-3, 1527S-1, 1527S-2, 1527(Bulk)
Basic UDI-DI	06082238401010000000005A4

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.

The Authorized European Representative for the concerned device(s) is

3M Deutschland GmbH
Health Care Business
Single Registration Number (TBD)
Carl-Schurz-Str. 1
41453 Neuss, Germany



Dianne Gibbs, Division Regulatory Affairs Manager
3M Company
2510 Conway Ave. St. Paul, MN 55144 USA

3 January 2020
Date

3M and Transpore are trademarks of 3M.