# *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 | Nexcare™ Transpore™ Clear Tape<br>Nexcare™ Foot Protection Tape<br>Spofaplast™ Transparent Tape<br>Viscoplast™ Foot Care Protective Tape<br>Nexcare™ Transparent Tape |
| Intended Purpose | A general-purpose tape used to secure most dressings, tubing and devices to skin. |
| Reference | NTT-1255, NTT-255, FPT-15, TAP1255TPOR, TAP255TPOR, V151NA1, N1540-1D |
| Basic UDI-DI | 06082238401050000000010DR |

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

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

Harry Janiski

Harry Janiski
Vice President of Quality
Medical Solutions Division
Health Care Business Group
3M Company

9 Jan 2023
Date

3M, Nexcare, Spofaplast and Viscoplast are trademarks of 3M.