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	Microfoam™ Surgical Tape
Intended Purpose	A general-purpose tape used to secure dressings in compression applications and devices to skin.
Catalogue Number	1528-1, 1528-2, 1528-3, 1528-4, 1528 (Bulk)
Basic UDI-DI	06082238401010000000004A2

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 DE-AR-000011642
Carl-Schurz-Str. 1
41453 Neuss, Germany

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

17 December 2021
Date

3M and Microfoam are trademarks of 3M.