

**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

Date

3M and Microfoam are trademarks of 3M.