EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Deutschland GmbH
Health Care Business
Carl-Schurz-Str. 1
41453 Neuss
Germany

Single Registration Number DE-MF-000011641

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

Trade Name	Micropore™
Intended Purpose	3M™ Micropore™ Surgical Tape is a general-purpose gentle tape used to secure dressings, lightweight tubing, and devices to skin.
Reference	1530-0, 1530-1, 1530-2, 1530-3 1530-0S, 1530-1S, 1530-2S, 1530-3S
Basic UDI-DI	06082327610100000000006CP

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.



November 29, 2022

Harald Ceschinski
Manager Regulatory Affairs and
Quality Management System
Health Care Business EMEA
3M Deutschland GmbH

Date

3M is a trademark of 3M.