EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

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

hereby declare under our sole responsibility that the following CE marked devices

Trade Name	Micropore™
Intended Purpose	Micropore™ Surgical Tape is a general-purpose gentle tape used to secure dressings, lightweight tubing, and devices to skin.
Reference	1530IP-1MD, 1530P-1SD, 1530P-1D, 1530P-1SD, 1530SP-1D, 1530SP-0D, 1530P-0D, 1530P-1D, 1535E-0, 1535E-1, 1535E-2, 1530IP-1S, 1530IP-4, 1530/1, 1530/5, 1530P-1S, 1530P-2S, 1530P-0S, 1530NP-1S, 1530NP-0SD, 1530NP-1SD, 1530SP-1, 1530IP-2S
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.

02. June 2020
Date

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