



## 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-OD, 1530P-OD, 1530P-1D, 1535E-0, 1535E-1, 1535E-2, 1530IP-1S, 1530IP-4, 1530/1, 1530/5, 1530P-1S, 1530P-2S, 1530P-OS, 1530NP-1S, 1530NP-OSD, 1530NP-1SD, 1530SP-1, 1530IP-2S
Basic UDI-DI	06082232761010000000006CP

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.

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

02. June 2020

Date

3M is a trademark of 3M.