EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Deutschland GmbH
Health Care Business
Single Registration Number DE-MF-000011641
Carl-Schurz-Str. 1
41453 Neuss
Germany

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

Trade Name	**3M™ Medipore™ + Pad**
Intended Purpose	**Adhesive Wound Dressing**
Reference	**3562E, 3564E, 3566E, 3569E, 3570E, 3571E, 3573E 3562NP, 3566NP, 3569NP, 3570NP, 3562SP, 3566SP, 3562P-10, 3566P-10, 3569P-10, 3570P-10, 3562PR-10,3566PR-10**
Basic UDI-DI	**06082327610100000000032CQ**

are classified per rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class 1 sterile devices in accordance with Annex IX and all other applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.

This declaration is made based on the quality assurance certificate
EU Certificate Number: 003626 MDR2017Q
Issued by: DQS Medizinprodukte GmbH, No. 0297

Claudia Inden
Regulatory Affairs Manager
3M Deutschland GmbH

November 11, 2024
Date