



Document Version: 1

## *Declaration of Conformity*

We

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

hereby declare under our sole responsibility  
that the CE marked products, to which this declaration relates,

**3M™ Medipore™ + Pad  
Adhesive Wound Dressing  
3562E, 3564E, 3566E, 3569E, 3570E, 3571E, 3572E, 3573E  
3562NP, 3566NP, 3569NP, 3570NP  
3562SP, 3566SP, 3569SP,  
3562P-10, 3566P-10, 3569P-10, 3570P-10  
3562P, 3566P, 3569P, 3570P  
3562IP-10, 3566IP-10, 3569IP-10, 3570IP-10**

are classified per rule 4 of Annex IX of the Medical Device Directive 93/42/EEC,  
as **Class I sterile** devices

and

are in accordance with  
*Annex II and all other applicable provisions of the Directive 93/42/EEC*  
on the approximation of the laws of the European Union Member States concerning medical devices.

This declaration is made on the basis of the quality assurance certificate  
EC Certificate No. 003626 MR2 delivered by DQS Medizinprodukte GmbH,  
August-Schanz-Straße 21, D-60433 Frankfurt am Main, No. 0297

Signature: \_\_\_\_\_

Margaret Bessenbach  
Manager Regulatory Affairs and Quality  
Health Care Business EMEA  
3M Deutschland GmbH

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May 11, 2020

Date