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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

May 11, 2020
Date

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