



3M Deutschland GmbH

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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™ Tegaderm™ I.V.
I.V. Transparent Film Dressing with Border
1633**

**3M™ Tegaderm™ I.V.
Transparent I.V. Film Dressing
1633P**

**3M™ Tegaderm™ Transparent
I.V. Transparent Film Dressing with Border
1635**

**3M™ Tegaderm™ Film
Transparent Film Dressing Frame Style
1623W**

are classified per rule 4 of Annex IX of the Medical Device Directive 93/42/EEC,
as **Class IIa** 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 valid until 2024-05-26 delivered by DQS Medizinprodukte GmbH,
August-Schanz-Straße 21, D-60433 Frankfurt am Main, No. 0297

Signature:

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

July 15, 2021

Neuss / Date

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