

**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	Tegaderm™ Transparent Film Dressing Tegaderm™ I.V. Transparent I.V. Film Dressing
Intended Purpose	Dressing can be used to cover and protect catheter sites and wounds, to maintain a moist environment for wound healing or to facilitate autolytic debridement, as a secondary dressing, as a protective cover over at-risk skin, to secure devices to the skin, to cover first and second degree burns and as a protective eye covering.
Reference (Products made in Germany)	1624P, 1624P-10, 1624NP, 1626P, 1626IP-10, 1626NP, 1626P-10, 1630NP, 1632P-10, 1633P
Basic UDI-DI (Products made in Germany)	0608223276101000000000CB
Reference (Products made in USA)	1622NP, 1630NP
Basic UDI-DI (Products made in USA)	06082232761010000000051CU

are classified per rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class 2a 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.

EC Certificate Number: 003626 MDR2017Q

Issued by: DQS Medizinprodukte GmbH, No. 0297

Claudia Inden
Senior Manager Regulatory Affairs
3M Deutschland GmbH

December 13, 2024

Date