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** IV Transparent Film Dressing with Border, **Tegaderm™ Film** Transparent Film Dressing Frame Style
Intended Purpose	IV Transparent Film Dressing with Border, Transparent Film Dressing Frame Style
Reference	**1633, 1635,** **1623W**
Basic UDI-DI	06082232761010000000000CB

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

Margaret Bessenbach

September 13, 2022

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

Date

Related to REG-STED-MDR-05-522836