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™ I.V. Advanced Securement Dressing**
Intended Purpose	Dressing can be used to cover and protect catheter insertion sites and minor wounds and to secure devices to the skin.
Reference (Products made in Germany)	**1681, 1683, 1685, 1688**
Basic UDI-DI (Products made in Germany)	06082327610100000000055D4
Reference (Products made in USA)	**1680, 1682, 1683, 1685, 1688, 1689, 1882**
Basic UDI-DI (Products made in USA)	06082327610100000000053CY

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

November 28, 2023

Harald Ceschinski
Senior Manager Regulatory Affairs and Quality
Health Care Business EMEA
3M Deutschland GmbH

Date

Related to REG-STED-MDR-05-709880