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

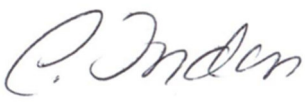
hereby declare under our sole responsibility that the following CE marked devices

Trade Name	Tegaderm™ Silicone Foam Border Dressing Tegaderm™ Silicone Foam Dressing
Intended Purpose	Tegaderm Silicone Foam Border Dressing and Tegaderm Silicone Foam Dressing are intended for management of low- to highly exuding partial and full thickness wounds
Reference	90640, 90641, 90642, 90643, 90646, 90647, 90648, 90631, 90632
Basic UDI-DI	06082232761010000000046D3 06082232761010000000047D5

are classified per rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class IIb 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 certificate
EU Quality Management Certificate: 003626MDR2017Q

Issued by: DQS Medizinprodukte GmbH, No. 0297
August-Schanz Straße 21, D-60433 Frankfurt am Main



Claudia Inden
Manager Regulatory Affairs
Medical Surgical Business
3M Deutschland GmbH

Neuss, June 10, 2024
Location/Date