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 device

Trade Name	Tegaderm™ High Performance Foam Adhesive Dressing
Intended Purpose	Tegaderm™ High Performance Foam Adhesive Dressing is indicated for use as a primary dressing for low - to highly exuding, partial and full thickness dermal wounds, including pressure ulcers, venous leg ulcers, abrasions, arterial ulcers, skin tears, neuropathic ulcers.
Reference	90610, 90611, 90612, 90613, 90614, 90616, 90619
Basic UDI-DI	06082327610100000000034CU

is 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 Management Certificate
EU Certificate Number: 003626 MDR2017Q
Issued by: DQS Medizinprodukte GmbH, August-Schanz Straße 21, D-60433 Frankfurt am Main, No. 0297

Margaret Bessenbach

Neuss, September 6, 2022

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

Location/Date