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	3M Tegaderm™ Roll
Intended Purpose	3M Tegaderm™ Roll Transparent Film Dressing is intended for use as a secondary dressing (e.g. used over and in combination with a primary sterile dressing); as a protective cover over at risk, intact skin; to secure devices to the skin; and as a waterproof fixation cover (e.g. to protect devices and primary dressings from outside fluid or water).
Reference	16002, 16004, 16006, 16004S
Basic UDI-DI	06082232761010000000016CS

are classified per rule 1 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class I devices in accordance with all applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.



February 6, 2025

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

Date

3M is a trademark of 3M.