EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Deutschland GmbH
Health Care Business
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	06082327610100000000016CS

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.

Margaret Bessenbach

September 09, 2020
Date

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