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 device*(s)*

Trade Name	Scotchcast™ Longuette
Intended Purpose	Scotchcast™ Longuettes are intended for use as a reinforcing strip for various types of lightweight synthetic casts, as well as specialised prosthetics and orthotic devices. 3M™ Scotchcast™ Longuettes can be used by itself to provide individual splinting or in combination with rigid or semi-rigid 3M™ Casting Tapes. Specific applications should be the responsibility of a qualified on site medical professional.
Reference	82199, 82299, 82203, 82200, 82201, 82202
Basic UDI-DI	06082327610100000000029D3

are classified per rule 1 Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class I non-sterile 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

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

25. August 2020
Date