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™ Plus
Intended Purpose	Scotchcast™ Plus is intended to construct rigid casts in order to immobilize fractures. 3M™ Scotchcast™ Plus tape can be used in the construction of most common orthopedic casts, as well as specialized prosthetics and orthotic devices. Suitability of the device for the particular application is the responsibility of a qualified, on-site medical professional.
Reference	82001, 82002, 82003, 82004, 82005, 82002L, 82003L, 82004L, 82002Y, 82003Y, 82002R, 82003R, 82004R, 82002B, 82003B, 82004B, 82005B, 82002G, 82003G, 82004G, 82002U, 82003U, 82004U, 82002X, 82003X, 82004X, 82002A, 82003A, 82004A, 82002V, 82003V, 82002W, 82003W
Basic UDI-DI	06082327610100000000031CN

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

14. July 2020
Date

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