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™ Quick Step Splint Double Sided Felt Splint
Intended Purpose	Scotchcast™ Quick Step Splint Double Sided Felt Splint is intended for use in the construction of common orthopedic/trauma splints. Specific splinting application suitability should be the responsibility of a qualified, on-site medical professional
Reference	75210, 75312, 75335, 75415, 75430, 75530
Basic UDI-DI	06082327610100000000025CT

are classified per rule 1 of 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.

09. February 2021
Date

Harald Ceschinski
Manager Regulatory Affairs and
Quality Management System
Health Care Business EMEA
3M Deutschland GmbH