EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Company
Single Registration Number (TBD)
2510 Conway Ave. St. Paul, MN 55144 USA

hereby declare under our sole responsibility that the following CE marked device(s)

Trade Name	Nexcare™ Blister Foot Care Plasters Nexcare™ Heel Blister Foot Care Plasters Nexcare™ Toe Blister Foot Care Plasters Viscoplast™ Hydrocolloid Blister Plasters Spofaplast® Hydrocolloid Blister Plasters
Intended Purpose	To cover and protect minor wounds
Reference	HBB-5N, N1406AS..2N, TBB-6N, V144NABN, V145NASN, 911N, 912N, 913N
Basic UDI-DI	06082238401050000000039EH

are classified per rule 4 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.

The Authorized European Representative for the concerned device(s) is

3M Deutschland GmbH
Health Care Business
Single Registration Number (TBD)
Carl-Schurz-Str. 1
41453 Neuss, Germany

10 DEC 2020

Bryan Becker, Division Quality and Regulatory Leader
3M Company
2510 Conway Ave. St. Paul, MN 55144 USA

Date