EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Company
Single Registration Number: US-MF-000014086
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 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	N1406AS_2N, HBB-5N, 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

EU Representative Address
3M Deutschland GmbH
Health Care Business
DE-AR-000011642
Carl-Schurz-Str. 1
41453 Neuss, Germany

Harry Janiski

Harry Janiski
Vice President of Quality
Medical Solutions Division
Health Care Business Group
3M Company

5 Dec 2022

Date

3M, Nexcare, Viscoplast, and Spofaplast are trademarks of 3M.