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	3M™ Nexcare™ Duo Bandages, 3M™ Spofaplast® Duo Bandages, 3M™ Viscoplast™ Duo Bandages
Intended Purpose	To cover and protect minor wounds
Reference	N17-20-3, N17-8-2, N17-5-1M, N17-40-4, N17-500-1, 613, 612, V17-20-3, N17-8-2, V17-3-1S, N17-3-1S
Basic UDI-DI	06082238401050000000045EC

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.

EU Authorized Representative:

EU Representative Address
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Bryan Becker, Senior Regulatory and Quality Manager
3M Consumer Health and Safety Division
2510 Conway Ave. St. Paul, MN 55144 USA

25FEB2022
Location/Date

3M is a trademark of 3M.