



EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Deutschland GmbH
Single Registration Number: DE-MF-000011641
Carl-Schurz-Str. 1
41453 Neuss
Germany

hereby declare under our sole responsibility that the following CE marked devices

Trade Name	3M™ Nexcare™ Bandages ultra stretch flexible comfort 3M™ Nexcare™ Soft'n flex 3M™ Nexcare™ Bandages Soft'n flex 360° natural feel bandages 3M™ Nexcare™ Bandages finger plaster flexible comfort
Intended Purpose	A general-purpose strips used to cover and protect minor wounds.
Reference	N1170B, N17030100, N1130ASD01W, 575-10E, SF-30D, SF-50D, NFP001W, N1130ASD03W, N1130ASD04
Basic UDI-DI	06082232761050000000010GA

are classified per rule 4 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.

27 November 2025

Wenny Sumarni
EMEA Regulatory Affairs Manager
3M Consumer Health Care

Date

3M is a trademark of 3M.