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 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.

April 16, 2020
Date

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