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 flexible foam active 3M™ Nexcare™ active 360° bandages 3M™ Nexcare™ Active™ strips
Intended Purpose	A general-purpose strips used to cover and protect minor wounds.
Reference	N1070B, N17022250, N17020100, N1030ASD01W, N1010ASDX02, N1010ASDX01, N17021100, N1030ASD03W, N1030ASD04, 576-50DP, 556-24DP, 572-30DP.
Basic UDI-DI	06082232761050000000009GR

are classified per rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class 1 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.

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

08. April 2020
Date