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™ ColdHot Therapy Pack 3M™ Nexcare™ ColdHot Therapy Pack, Back & Abdomen Belt 3M™ Futuro™ Hot/Cold Therapy Pack 3M™ Futuro™ ColdHot Compress
Intended Purpose	The reusable 3M™ Nexcare™ ColdHot Therapy Pack Mask is intended to provide a superficial cold or heat to affected body parts.
Reference	N1573DAB, N1573ECH, N1573B, N1570, N1570B, N1578DAB, N1578B, N1570B2, N15710IE, N1573G, N1570G, N1578G, N1571TI-DAB, N1573KID, 02070, F1570, N15711S, N15711L.
Basic UDI-DI	06082232761050000000014GJ

are classified per rule 1 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class 1 devices in accordance with all applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.

31. March 2020
Date

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