# *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<br>3M™ Nexcare™ ColdHot Therapy Pack, Back & Abdomen Belt<br>3M™ Futuro™ Hot/Cold Therapy Pack<br>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