



EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Deutschland GmbH
Carl-Schurz-Str. 1
41453 Neuss
Germany

Single Registration Number: DE-MF-000011641

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.

25 July 2025

Wenny Sumarni
EMEA Regulatory Affairs Manager
3M Consumer Health Care

Date

3M is a trademark of 3M.