Confidential

Declaration of Conformity

As Legal Manufacturer, we

3M Company
Single Registration Number US-MF-000014086
2510 Conway Ave. St. Paul, MN 55144 USA

hereby declare under our sole responsibility that the following CE marked device(s)

Trade Name*	Coban™ Non-Latex Self-Adherent Wrap, (non-sterile)
Intended Purpose	Elastic wrap used to provide compression, support or to secure dressings or devices.
Reference	2081, 2081B, 2081O, 2081P, 2081T, 20815, 20815B, 2082, 2082B, 2082G, 2082N, 2082O, 2082P, 2083, 2083B, 2083O, 2083P, 2084, 2084B, 2084O, 2084P, 2084L, 2086, 2081C, 20815C, 2082C, 2083C, 2084C, 2082-1X, 2083-1X, 2084-1X, 2086-1X, 2084K (Bulk), 2084LK (Bulk), 2086K (Bulk)
Basic UDI-DI	06082238401010000000109V

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

The Authorized European Representative for the concerned device(s) is

3M Deutschland GmbH
Health Care Business
Single Registration Number DE-AR-000011642
Carl-Schurz-Str. 1
41453 Neuss, Germany

Dianne Gibbs

23 February 2022

Dianne Gibbs, RAC
Regulatory Affairs Director
3M Medical Solutions Division

Location/Date

3M and Coban are trademarks of 3M.