EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Company
Single Registration Number *(TBD)*
2510 Conway Ave. St. Paul, MN 55144 USA

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

Trade Name*	Coban™ Self-Adherent Wrap (non-sterile)
Intended Purpose	Elastic wrap used to provide compression, support or to secure dressings or devices.
Reference	2081, 20815, 2082, 2083, 2084, 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 (TBD)
Carl-Schurz-Str. 1
41453 Neuss, Germany

Dianne Gibbs, Division Regulatory Affairs Manager
3M Company
2510 Conway Ave. St. Paul, MN 55144 USA

21 May 2020
Date

3M and Coban are trademarks of 3M.

Issued to Authorized Representative