

## Declaration of Conformity

*As Legal Manufacturer, we*

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

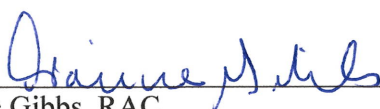
*hereby declare under our sole responsibility that the following CE marked device(s)*

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	060822384010100000000109V

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, RAC  
Regulatory Affairs Director  
3M Medical Solutions Division

*23 February 2022*  
Location/Date

3M and Coban are trademarks of 3M.

**3M™ Coban™ NL Self-Adherent Wrap (non-sterile) – Declaration of Conformity (MDR)**

*NOTE: Paper Copies are Uncontrolled unless Stamped 'Controlled Copy' in Red*

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