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	Coban™2Lite
Intended Purpose	Coban 2 Lite Compression System may be used to provide compression for various medical diagnoses, such as: lymphedema and venous leg ulcers. Coban 2 Lite Compression System is recommended for use in patients with an ABPI > 0.5 or for those limbs with small circumferences. It is used mainly for the arms, fingers, toes and lower extremities of patients with mixed arterio-venous disease or poor tolerance of high compression.
Reference	2794E, 20751, 20721,20723, 20724, 20726, 20713, 20714, 20716
Basic UDI-DI	06082327610100000000004CK

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.

Margaret Bessenbach

Margaret Bessenbach
Manager Regulatory Affairs and Quality
Health Care Business EMEA
3M Deutschland GmbH
3M is a trademark of 3M.

March 17, 2020
Date

Relating to REG-STED-MDR-05-630644