

**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	Cavilon™ Durable Barrier Cream
Intended Purpose	Cream for protection of intact and injured skin from damage due to bodily fluids or to moisturize and condition dry skin.
Reference	3391G: 28g (1oz) tube 3392G: 92g (3.5oz) tube 3392GS: 2g sachet
Basic UDI-DI	06082238401010000000017AB

are classified per rules 1 and 4 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 devices 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

13 May 2020  
Date

3M and Cavilon are trademarks of 3M.