EUROPEAN MEDICAL DEVICE REGULATION

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	3M™ Cavilon™ Advanced Skin Protectant
Intended Purpose	Polymeric solution intended to cover and protect intact or damaged skin.
Reference	5050G, 5050G4P: 2.7 mL solution 5051G, 5051G4P: 0.7 mL solution
Basic UDI-DI	06082238401010000000064AL

are classified per Rule 4 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class IIa devices in accordance with Annex XI Part A and all other applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.

This declaration is made based on the Quality Assurance Certificate
EC Certificate Number: MDR 725202
Issued by: BSI, 2797

EU Authorized Representative:

EU Representative Address
3M Deutschland GmbH
Health Care Business
DE-AR-000011642
Carl-Schurz-Str. 1
41453 Neuss, Germany

DocuSigned by:
Mary Fretland
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Mary Fretland
Manager, Regulatory Affairs
3M Medical Surgical Division

8/2/2024
Date