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 devices

Trade Name	1. 3M™ Single use Pivoting Clipper Blade Assembly 2. 3M™ Single use Specialty Clipper Blade Assembly
Intended Purpose	Surgical Clipper Blade
Reference	1. 9660 2. 9690
Basic UDI-DI	06082238401010000000051AB

are classified per rule 13 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.

EU Authorized Representative:

3M Deutschland GmbH
Health Care Business
Single Registration Number: DE-AR-000011642
Carl-Schurz-Str. 1
41453 Neuss, Germany

DocuSigned by:
Nadia Battah
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Nadia Battah
Regulatory Affairs Manager
3M Company
Medical Solutions Division

2/27/2023
Date