

**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

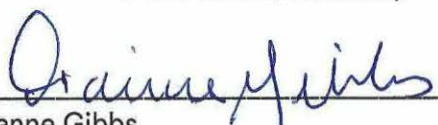
hereby declare under our sole responsibility that the following CE marked devices

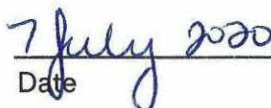
Trade Name	3M™ Single use Pivoting Clipper Blade Assembly and 3M™ Single use Specialty Clipper Blade Assembly
Intended Purpose	Surgical Clipper Blade
Reference	9660 and 9690
Basic UDI-DI	06082238401010000000051AB

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.

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


Date

Issued to Authorized Representative (EC REP)

3M is a trademark of 3M.