# *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

7 July 2020
Date

*Issued to Authorized Representative (EC REP)*



**THIS DOCUMENT WAS PRINTED ON 29-Jun-2020 8:50 PM**