EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we
3M Deutschland GmbH
Health Care Business
Carl-Schurz-Str. 1
41453 Neuss
Germany

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

Trade Name	Surgical Masks
Intended Purpose	The intended use of 1810F and 1810G surgical masks is to protect patients from microbial dissemination of infective agents from the wearer.
Reference	1810F, 1810G
Basic UDI-DI	06082327610100000000010CE

are classified per rule 1 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.

Margaret Bessenbach

May 27, 2020
Date

Margaret Bessenbach
Manager Regulatory Affairs and Quality
Health Care Business EMEA
3M Deutschland GmbH