EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Deutschland GmbH
Health Care Business
Single Registration Number: DE-MF-000012859
Carl-Schurz-Str. 1
41453 Neuss
Germany

hereby declare under our sole responsibility that the following CE marked device(s)

Trade Name	Protemp 4
Intended Purpose	temporary restorative material for temporary restoration on prepared teeth
Reference	46953, 46954, 46956, 46957, 46959, 46960, 46972
Basic UDI-DI	06082327610200000000040DN

is/are classified per rules 8 and 19 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class IIa device(s) in accordance with Annex IX and all other applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.

We hereby declare under our sole responsibility that the following CE marked accessory(ies) which are intended to be used together with the above medical device(s)

Accessory(ies)	Reference	Basic UDI-DI	Rule(s) of Annex VIII	Class
Garant Dispenser 10:1	77581	06082232761020000000015DP	1	I
Mixing Tips blue	71453	06082232761020000000016DR	5	I

is/are classified according to Annex VIII of the Medical Device Regulation (EU) 2017/745 in accordance with all 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
EU QMS Certificate (MDR): G10 078535 0040
Issued by: TÜV SÜD Product Service GmbH (identification no. 0123)

Dr. Desi W. Soegiarto
Manager Regulatory Medical Devices
3M Deutschland GmbH

Seefeld/December 13, 2022
Location/Date