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EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Company, ESPE Dental Products
Single Registration Number: US-MF-000014051
2510 Conway Ave. St. Paul, MN 55144 USA

hereby declare under our sole responsibility that the following CE-marked device

Trade Name	3M™ Filtek™ Easy Match Universal Restorative
Intended Purpose	Dental composite resin for anterior and posterior restorations
Reference	6210B| 6210N| 6210W| 6210K| 6210S| 6210| 6220B| 6220N| 6220W| 6220K| 6220S| 6220
Basic UDI-DI	06082238401020000000047BK

is classified per rule 8 and 19 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class IIa devices 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.

This declaration is made based on the EU Quality Assurance Certificate G10 117189 0001 Issued by TUV SUD Product Service GmbH (identification no. 0123)

EU Authorized Representative:

EU Representative Address
3M Deutschland GmbH
Health Care Business
Single Registration Number:
DE-AR-000012860
Carl-Schurz-Str. 1
41453 Neuss, Germany

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DocuSigned by:
Kristin Totushek
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Kristin Totushek
Senior Regulatory Affairs Manager
3M Oral Care Solutions Division

4/23/2024 (St Paul, MN)
Location/Date

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