Declaration of Conformity

As Legal Manufacturer, we

3M ESPE Dental Products
2510 Conway Ave. St. Paul, MN 55144 USA

hereby declare under our sole responsibility
that the following UKCA marked products to which this declaration relates;

3M™ Clinpro™ Clear Fluoride Treatment
Product Reference Numbers: 7300FF, 7300M, 7300W, 7310FF, 7310M, 7310M, 7320SP

are classified, per Annex IX of Council Directive 93/42/EEC
as implemented in the UK through the Medical Devices Regulations 2002 (SI618)
as subsequently amended by the EU Exit Regulations of 2019 (SI 791) and 2020 (SI 1478).
as Class I Medical Devices

and meet the relevant essential requirements, of Annex I of Council Directive 93/42/EEC
as implemented in the UK through the Medical Devices Regulations 2002 (SI618)
as subsequently amended by the EU Exit Regulations of 2019 (SI 791) and 2020 (SI 1478).

UK Responsible Person:
3M United Kingdom PLC
3M Centre, Cain Road,
Bracknell, RG12 8HT,
United Kingdom.

DocuSigned by:
Katelyn Manning
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Katelyn Manning, RAC
Sr. Regulatory Affairs Manager
3M Oral Care Solutions Division

5/20/2024
Date