Declaration of Conformity

As Legal Manufacturer, we,
3M Company,
2510 Conway Ave
St Paul, Minnesota 55144
USA

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

3M™ Medipore™ H Soft Cloth Surgical Tape
Product Reference Numbers: 2861, 2862, 2863, 2864, 2866, 2868, 2862S, 2864S, 2866S, 2860S-1, 2860S-2, 2860S-4, 2860S-6, 2860-6, 2860S-2U, 2860S-4U, 2860S-6U

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.

Dianne Gibbs

Dianne Gibbs, RAC
Regulatory Affairs Director
3M Medical Solutions Division

11 July 2022
Date