



## ***Declaration of Conformity***

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

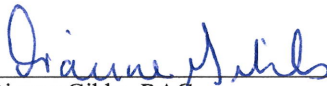
hereby declare under our sole responsibility  
that the following UKCA marked products to which this declaration relates;

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.

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

  
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Date