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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:

1. 3M™ Littmann® Cardiology IV™ Stethoscopes
2. 3M™ Littmann® Classic III™ Stethoscopes
3. 3M™ Littmann® Master Cardiology™ Stethoscopes
4. 3M™ Littmann® Classic II Infant Stethoscopes
5. 3M™ Littmann® Classic II Pediatric Stethoscopes
6. 3M™ Littmann® Lightweight II S.E. Stethoscopes

Product Reference Numbers:
1. Catalog Numbers: 6151, 6152, 6154, 6155, 6156, 6158, 6159, 6163, 6165, 6168, 6170, 6176, 6177, 6179, 6184, 6190, 6200, 6201, 6202, 6203, 6204, 6205, 6232, 6234, 6238, 6239, 6240, 6241, 6242.
2. Catalog numbers: 5620, 5621, 5622, 5623, 5627, 5630, 5633, 5648, 5803, 5806, 5807, 5809, 5811, 5831, 5832, 5835, 5839, 5861, 5863, 5864, 5868, 5870, 5872, 5873, 5875, 5959, 5960, 5962.
3. Catalog Numbers: 2160, 2161, 2163, 2164, 2167, 2175, 2176.
4. Catalog Numbers: 2114, 2114R, 2124, 2157.
5. Catalog Numbers: 2113, 2113R, 2119, 2122, 2153.
6. Catalog Numbers: 2450, 2451, 2452, 2454, 2456.

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:
Nadia Battah
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Nadia Battah
Regulatory Affairs Manager
3M Company,
2510 Conway Ave
St Paul, Minnesota 55144 USA

8/21/2023
Date