



EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Company
Single Registration Number (TBD)
2510 Conway Ave. St. Paul, MN 55144 USA

hereby declare under our sole responsibility that the following CE marked devices

| | |
|------------------|--|
| Trade Name | <ol style="list-style-type: none">1. Littmann® Cardiology IV™ Stethoscope2. Littmann® Classic III™ Stethoscope3. Littmann® Classic II Pediatric Stethoscope4. Littmann® Master Cardiology™ Stethoscope5. Littmann® Master Classic II™ Stethoscope6. Littmann® Classic II SE Stethoscope7. Littmann® Classic II Infant Stethoscope8. Littmann® Lightweight II SE Stethoscope |
| Accessories | None. |
| Intended Purpose | Stethoscope (mechanical) |
| Reference | <ol style="list-style-type: none">1. 6151, 6152, 6154, 6155, 6156, 6158, 6159, 6162, 6163, 6164, 6165, 6166, 6168, 6170, 6171, 6176, 6177, 6179, 6180, 6181, 6182, 6183, 6184, 6190, 6200, 6201, 6202, 6203, 6204, 6205, 6206, 6232, 6234, 6238, 6239, 6240, 6241, 62422. 5620, 5621, 5622, 5623, 5627, 5630, 5633, 5646, 5647, 5648, 5803, 5806, 5807, 5809, 5811, 5812, 5831, 5832, 5835, 5839, 5861, 5862, 5863, 5864, 5868, 5870, 5871, 5872, 5873, 5874, 5875, 5959, 5960, 59623. 2113, 2113R, 2119, 2122, 21534. 2160, 2161, 2163, 2164, 2167, 2175, 2176, 2178, 21825. 1392, 2141, 2144L, 2146, 21476. 21387. 2114, 2114R, 2124, 21578. 2450, 2451, 2452, 2454, 2456 |
| Basic UDI-DI | <ol style="list-style-type: none">1. 06082238401010000000026AC2. 06082238401010000000027AE3. 06082238401010000000028AG4. 06082238401010000000029AJ5. 06082238401010000000030A3 |



| | |
|--|------------------------------|
| | 6. 06082238401010000000031A5 |
| | 7. 06082238401010000000032A7 |
| | 8. 06082238401010000000033A9 |

are classified per rule 1 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class I devices in accordance with all applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.

The Authorized European Representative for the concerned devices is

3M Deutschland GmbH
Health Care Business
Single Registration Number (TBD)
Carl-Schurz-Str. 1
41453 Neuss, Germany

Dianne Gibbs
Division Regulatory Affairs Manager
3M Company
2510 Conway Ave. St. Paul, MN 55144 USA

18 February 2021

Date

3M, Littmann, Cardiology IV, Classic III, Master Cardiology, and Master Classic II are marks and/or registered marks of 3M.