



## 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"><li>1. Littmann® Cardiology IV™ Stethoscope</li><li>2. Littmann® Classic III™ Stethoscope</li><li>3. Littmann® Classic II Pediatric Stethoscope</li><li>4. Littmann® Master Cardiology™ Stethoscope</li><li>5. Littmann® Master Classic II™ Stethoscope</li><li>6. Littmann® Classic II SE Stethoscope</li><li>7. Littmann® Classic II Infant Stethoscope</li><li>8. Littmann® Lightweight II SE Stethoscope</li></ol>
Intended Purpose	Stethoscope
Reference	<ol style="list-style-type: none"><li>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, 6240, 6241, 6242</li><li>2. 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, 5962</li><li>3. 2113, 2113R, 2119, 2122, 2153</li><li>4. 2160, 2161, 2163, 2164, 2167, 2175, 2176, 2178, 2182</li><li>5. 1392, 2141, 2144L, 2146, 2147</li><li>6. 2138</li><li>7. 2114, 2114R, 2124, 2157</li><li>8. 2450, 2451, 2452, 2454, 2456</li></ol>
Basic UDI-DI	<ol style="list-style-type: none"><li>1. 06082238401010000000026AC</li><li>2. 06082238401010000000027AE</li><li>3. 06082238401010000000028AG</li><li>4. 06082238401010000000029AJ</li></ol>



	5. 06082238401010000000030A3 6. 06082238401010000000031A5 7. 06082238401010000000032A7 8. 06082238401010000000033A9
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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)  
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Dianne Gibbs  
Division Regulatory Affairs Manager  
3M Company  
2510 Conway Ave. St. Paul, MN 55144 USA

7 July 2020  
Date

3M is a trademark of 3M.