



## 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>
Accessories	None.
Intended Purpose	Stethoscope (mechanical)
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, 6239, 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><li>5. 06082238401010000000030A3</li></ol>



	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

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

18 February 2021

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Date

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