3M Health Care Business

3M Center
2510 Conway Ave, Bldg. 275-5W-06
St. Paul, MN 55144 U.S.A.
651 733 1110

Declaration of Conformity

As Legal Manufacturer
We, 3M Company, 3M Health Care,
3M Center, 2510 Conway Ave, Bldg. 275-5W-06
Saint Paul, MN 55144 USA
hereby declare under our sole responsibility
that the CE marked products to which this declaration relates,

Product numbers:

3M™ Littmann® Traditional Stethoscope	3141, 3142, 3143
3M™ Littmann® Master Cardiology™ Stethoscope	2159, 2160, 2161, 2163, 2164, 2165, 2167, 2175, 2176, 2178, 2182, 2183
3M™ Littmann® Cardiology STC Stethoscope	4471, 4472, 4473, 4474, 4475
3M™ Littmann® Cardiology III™ Stethoscope	3127, 3128, 3128BRS, 3129, 3130, 3131BE, 3134, 3135, 3136, 3137, 3137CPR, 3138, 3140, 3146, 3148, 3149, 3152RBW, 3157SM, 3158, 3159, 3161, 3163, 3164, 3165, 3166
3M™ Littmann® Cardiology IV™ Stethoscope	6151, 6152, 6153, 6154, 6155, 6156, 6157, 6158, 6159, 6161,6162, 6163, 6164, 6165, 6166, 6167, 6168, 6169, 6170, 6171, 6172, 6173, 6174, 6175, 6176, 6177, 6178, 6179
3M™ Littmann® Master Classic II™ Stethoscope	2139, 2141, 2142G, 2143, 2144L, 2146, 2147, 2630, 2632, 2633, 2634, 2636
3M™ Littmann® Classic III Stethoscope	5620, 5621, 5622, 5623, 5624, 5626, 5627, 5629, 5630, 5631, 5633, 5639, 5641, 5803, 5806, 5807, 5809, 5811, 5812, 5813, 5829, 5831, 5832, 5835, 5839, 5840, 5867, 5868, 5869, 5870, 5871
3M™ Littmann® Classic II S.E. Stethoscope	2138, 2201, 2201BRS, 2203, 2205, 2206, 2208, 2209, 2210, 2211, 2215, 2218BE, 2812, 2813, 2814, 2815, 2816, 2817, 2818, 2819, 2820CPR, 2822, 2823, 2827SM, 2828, 2829RBW, 2830, 2836, 2837, 2941, 2942
3M™ Littmann® Classic II Pediatric Stethoscope	2113, 2113R, 2115, 2119, 2122, 2123, 2131, 2136, 2153, 2154, 2155
3M™ Littmann® Classic II Infant Stethoscope	2114, 2114R, 2120, 2124, 2125, 2126, 2132, 2156, 2157, 2158, 2179
3M™ Littmann® Select Stethoscope	2290, 2291, 2292, 2293, 2294, 2296, 2298, 2301, 2303, 2305, 2306, 2310
3M™ Littmann® Lightweight II S.E. Stethoscope	2450, 2451, 2452, 2453, 2454, 2455, 2456

are classified,
per Rule 1 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC
as a Class I device, and

are in accordance with Annex VII of Directive 93/42/EEC, as amended per 2007/47/EC,
on the approximation of the laws of the European Member States concerning medical devices.

EU Representative Address
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Signature: Dianne Gibbs Date: 28 March 2017

Dianne Gibbs
3M Health Care
Division Regulatory Affairs Manager
Infection Prevention Division

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