



**Declaration of Conformity**

As Legal Manufacturer  
We, 3M Health Care Business,  
2510 Conway Ave  
St. Paul, MN 55144 USA

hereby declare under our sole responsibility  
that the CE marked products to which this declaration relates,

Littmann Traditional Stethoscope	3141, 3142, 3143
Littmann Master Cardiology	2159, 2160, 2161, 2163, 2164, 2165, 2167, 2168, 2169, 2175, 2176, 2178
Littmann Cardiology STC	4471, 4472, 4473, 4474, 4475
Littmann Cardiology III	3127, 3128, 3128BRS, 3129, 3130, 3131BE, 3134, 3135, 3136, 3137, 3137CPR, 3138, 3140, 3146, 3148, 3149, 3152RBW, 3157SM
Littmann Master Classic II	2139, 2141, 2142G, 2143, 2144L, 2146, 2147, 2630, 2632, 2633, 2634
Littmann Classic II S.E.	2138, 2201, 2201BRS, 2203, 2205, 2206, 2208, 2209, 2210, 2211, 2215, 2218BE, 2812, 2813, 2814, 2816, 2817, 2818, 2819, 2820CPR, 2822, 2823, 2815, 2827SM, 2829RBW
Littmann Classic II Pediatric	2113, 2113R, 2115, 2119, 2122, 2123, 2131, 2136, 2153, 2154, 2155
Littmann Classic II Infant	2114, 2114R, 2120, 2124, 2125, 2126, 2132, 2156, 2157, 2158, 2179
Littmann Select	2290, 2291, 2292, 2293, 2294, 2296, 2298, 2301, 2303, 2305, 2306, 2310
Littmann Lightweight II S.E.	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.

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