



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,

3M™ Littmann™ Traditional Stethoscope™	3141, 3142, 3143
3M™ Littmann™ Master Cardiology™	2159, 2160, 2161, 2163, 2164, 2165, 2167, 2175, 2176, 2178, 2182, 2183
3M™ Littmann™ Cardiology STC™	4471, 4472, 4473, 4474, 4475
3M™ Littmann™ Cardiology III™	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™ Master Classic II™	2139, 2141, 2142G, 2143, 2144L, 2146, 2147, 2630, 2632, 2633, 2634, 2636
3M™ Littmann™ Classic II S.E™	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™	2113, 2113R, 2115, 2119, 2122, 2123, 2131, 2136, 2153, 2154, 2155
3M™ Littmann™ Classic II Infant™	2114, 2114R, 2120, 2124, 2125, 2126, 2132, 2156, 2157, 2158, 2179
3M™ Littmann™ Select™	2290, 2291, 2292, 2293, 2294, 2296, 2298, 2301, 2303, 2305, 2306, 2310
3M™ 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.

EU Representative Address
3M Deutschland GmbH
Health Care Business
Carl-Schurz-Str. 1
41453 Neuss, Germany

Signature: Suzanne M. Danielson
Suzanne M. Danielson
3M Health Care
Vice President, Regulatory Affairs and Quality Assurance
Infection Prevention Division

Date: January 31, 2014