3M Health Care Business

3M Center
St. Paul, MN 55144-1000
651 733 1110

3M

Declaration of Conformity

We, 3M Health Care,

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

Littmann Master Cardiology	2159, 2160, 2161, 2163, 2164, 2165, 2167, 2168, 2169, 2178
Littmann Cardiology Soft Touch Chestpiece	4470, 4471, 4472, 4473, 4474, 4475, 4476, 4477
Littmann Cardiology III	3127, 3128, 3129, 3130, 3134, 3135, 3136, 3137
Littmann Cardiology III Black Edition	3131BE
Littmann Cardiology Brass Edition	2175
Littmann Cardiology Smoke Edition	2176
Littmann Master Classic II	2141, 2142G, 2143, 2144L, 2146, 2147, 2630, 2632, 2633, 2634
Littmann Classic II S.E.	2201, 2203, 2205, 2206, 2208, 2209, 2210, 2211, 2215, 2812, 2813, 2814, 2815, 2816, 2817, 2818,2819, 2820, 2821, 2822, 2823
Littmann Classic II S.E. Black Edition	2218BE
Littmann Classic II Pediatric	2113, 2113R, 2115, 2119, 2122, 2123, 2131, 2153
Littmann Classic II Infant	2114, 2114R, 2120, 2124, 2125, 2126, 2132, 2157
Littmann Select	2290, 2291,2292, 2293, 2294, 2296, 2297, 2298, 2301, 2302, 2303, 2305, 2306, 2307, 2309, 2310
Littmann Master Classic II Teaching	2139
Littmann Classic II S.E. Teaching	2138
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.

This certificate is valid for devices originating from the following sites:

3M Health Care
3M Brookings
601 22nd. Ave. South
Brookings, South Dakota USA 57006

EU Representative Address
3M Medica
Zweigniederlassung der 3M Deutschland GmbH
Trading as "3M Health Care"
Hammfelddamm 11 D-41453 Neuss, Germany

Signature: Suzanne M. Danielson Date: November 11, 2010

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

REG-DOC-05-132620, Littmann Mechanical Stethoscopes, Version 2,
Status: Release, Release Date: 11/9/2010 9:41:57 AM

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3M Health Care Business

3M Center
St. Paul, MN 55144-1000
651 733 1110

3M

Declaration of Conformity

We, 3M Health Care,

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

3M™ Littmann® Electronic Stethoscope Models
3100, 3200

is classified as a Class IIa active device,
according to Rule 10 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC

and

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

In addition, we declare that the above mentioned devices fulfil the applicable provisions of the Directive 93/42/EEC, as amended per 2007/47/EC.

This declaration is made on the basis of the EC certificate CE 00493 delivered by BSI, 0086

This certificate is valid for devices originating from the following sites:

Bang & Olufsen Medicom A/S (B&O)
Gimsinglundvej 20
DK-7600 Struer, Denmark

EU Representative Address
3M Medica
Zweigniederlassung der 3M Deutschland GmbH
Trading as "3M Health Care"
Hammfeldamm 11
D-41453 Neuss, Germany

Signature: _____ Date: 18 March 2010

Suzanne M. Danielson
3M Health Care
Regulatory Affairs and Quality Assurance
Infection Prevention Division

REG-DOC-05-000025, Model 3100 and 3200 Electronic Stethoscope, Version 5,
Status: Release, Release Date: 3/18/2010 11:41:58 AM