



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
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 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, 2818, 2829, 2820, 2821, 2822
Littmann Classic II S.E. Black Edition	2218BE
Littmann Classic II Pediatric	2113, 2113R, 2115, 2119, 2122, 2123, 2131
Littmann Classic II Infant	2114, 2114R, 2120, 2124, 2125, 2126, 2132
Littmann Select	2290, 2291, 2292, 2293, 2294, 2296, 2297, 2298, 2301, 2302, 2303, 2305, 2306, 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 a Class I device,
and

are in accordance with Annex VII of Directive 93/42/EEC
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

Signature:


Suzanne M. Danielson
Regulatory Affairs and Quality Director
Medical Division

Date:

