



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

Littmann Cardiology Soft Touch Chestpiece 4470, 4471, 4472, 4473, 4474 & 4475

Littmann Cardiology III 3127, 3128, 3129, 3130, 3134 & 3135

Littmann Master Classic II 2141, 2142G, 2143, 2144L, 2146, 2147, 2630, 2632 & 2633

Littmann Classic II S.E. 2201, 2203, 2205, 2206, 2208, 2209, 2210, 2211, 2215, 2812, 2813 & 2814

Littmann Classic II Pediatric 2113, 2113R, 2115, 2119, 2122 & 2123

Littmann Classic II Infant 2114, 2114R, 2120, 2124, 2125 & 2126

Littmann Select 2290, 2291, 2292, 2293, 2294, 2296, 2297, 2298, 2301, 2302 & 2303

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,

according to the rule of Annex IX of the Medical Device Directive 93/42/EEC,
as a Class I device,

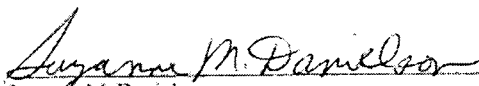
and

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

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