3M Company

2510 Conway Ave,
St. Paul, MN 55144 U.S.A.
651 733 1110

Declaration of Conformity

As Legal Manufacturer
We, 3M Company,
2510 Conway Ave,
St. Paul, MN 55144 USA
hereby declare under our sole responsibility
that the CE marked products to which this declaration relates,

Product numbers:

Littmann® Master Cardiology ™ Stethoscope	2160, 2161, 2163, 2164, 2167, 2175, 2176, 2178, 2182
Littmann® Cardiology IV ™ Stethoscope	6151, 6152, 6154, 6155, 6156, 6158, 6159, 6162, 6163, 6164, 6165, 6166, 6168, 6170, 6171, 6176, 6179, 6180, 6181, 6190, 6200, 6201, 6202, 6203, 6204, 6205, 6206, 6232, 6234, 6238, 6240, 6241, 6242
Littmann® Master Classic II™ Stethoscope	2141, 2144L, 2146, 2147, 1392
Littmann® Classic III ™ Stethoscope	5620, 5621, 5622, 5623, 5627, 5630, 5633, 5646, 5647, 5803, 5806, 5807, 5809, 5811, 5812, 5831, 5832, 5835, 5839, 5861, 5862, 5863, 5864, 5868, 5870, 5871, 5872, 5873, 5874, 5875, 5959, 5960, 5962
Littmann® Classic II S.E. Stethoscope	2138
Littmann® Classic II Pediatric Stethoscope	2113, 2113R, 2119, 2122, 2153,
Littmann® Classic II Infant Stethoscope	2114, 2114R, 2124, 2157,
Littmann® Lightweight II S.E. Stethoscope	2450, 2451, 2452, 2454, 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: Dianne L. Gibbs Date: 13 March 2020

Dianne L. Gibbs
3M Company
Division Regulatory Affairs Manager
Medical Solutions Division

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THIS DOCUMENT WAS PRINTED ON 13-Mar-2020 12:05 PM