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
Littmann Cardiology Soft Touch Chestpiece
Littmann Cardiology III
Littmann Cardiology III Black Edition
Littmann Cardiology Brass Edition
Littmann Cardiology Smoke Edition
Littmann Master Classic II
Littmann Classic II S.E.
Littmann Classic II S.E. Black Edition
Littmann Classic II Pediatric
Littmann Classic II Infant
Littmann Select
Littmann Master Classic II Teaching
Littmann Classic II S.E. Teaching
Littmann Lightweight II S.E.

are classified,

per Rule I of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC as a Class I device, and


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

Date: November 11, 2010

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REG-DOC-05-132620, Littmann Mechanical Stethoscopes, Version 2,

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

Date: 18 May 2010