# EC Declaration of Conformity

We hereby declare that the products of the product category

**Product name:**

**MDF 727E Singularis™ SOLO™ Stethoscope**
**MDF 747E Singularis™ DUET™ Stethoscope**
**MDF 787E Singularis™ VIVO™ Stethoscope**
**MDF 727 Single Head Stethoscope**
**MDF 727C Single Head Stethoscope**
**MDF 740 Pulse Time™ Stethoscope**
**MDF 740C Pulse Time™ Stethoscope**
**MDF 747 Dual Head Stethoscope**
**MDF 747C Dual Head Stethoscope**
**MDF 747XP Acoustica™ Stethoscope**
**MDF 757PT Pulse Time™ Teaching Stethoscope**
**MDF 767 Sprague Rappaport Stethoscope**
**MDF 767K Sprague Rappaport Stethoscope > 22k Gold**
**MDF 767X Deluxe Sprague Rappaport X Stethoscope**
**MDF 767XK Deluxe Sprague Rappaport X Stethoscope > 22K Gold**
**MDF 777 MD One™ Stainless Steel Dual Head Stethoscope**
**MDF 777C MD One™ Stainless Steel Dual Head Stethoscope**
**MDF 777I MD One™ Stainless Steel Dual Head Stethoscope**
**MDF 777K MD One™ Stainless Steel Dual Head Stethoscope > 22K Gold**
**MDF 787 Infant & Neonatal Stethoscope**
**MDF 787XP Deluxe Infant & Neonatal Stethoscope**
**MDF 797 Classic Cardiology™ Stethoscope**
**MDF 797K Classic Cardiology™ Stethoscope > 22K Gold**
**MDF 797DD ER Premier™ Stethoscope**
**MDF 797DDK ER Premier™ Stethoscope > 22K Gold**
**MDF 797X ProCardial X™ Stethoscope**
**MDF 797CC ProCardial™ C3 > Critical Cardiac Care Edition Stethoscope**

**Product Category (UMDNS-Code):** **13-750** Stethoscopes

Manufactured by **MDF Instruments Medifriend Inc.**
**3F Building 6, 1898 Lai Yin Road Jiu Ting Town,**
**Song Jiang District 201615 Shanghai China**

Fulfils the Essential Requirements of Annex I of the Directive 93/42/EEC and are manufactured and placed on the market under the sole responsibility of the manufacturer following the regulations of this directive.

The products are classified according to **Annex IX, rule I of MDD 93/42/EEC** as a medical device **class I**

Conformity Assessment Procedure: **MDD 93/42/EEC Annex VII**

| MDF Instruments<br>Medifriend Inc. | Declaration of Conformity<br>According<br>MDD 93/42/EEC | Page 2 of 2<br>Revision 03<br>Valid from Dec. 12th, 2013 |
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No Notified Body is used in Conformity Assessment for the above products

Authorised Representative in EU:

MT Promedt Consulting GmbH
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Shanghai Dec. 12, 2013

*Place, date*

*Legally binding signature, Function*