AMERICAN DIAGNOSTIC CORPORATION

Declaration of Conformity

We

declare with sole responsibility, that our product(s): *UMDNS code 13755 Stethoscopes, Mechanical of the following model(s) 5079-73 (672BKWA), 5079-74 (672RWA), 5079-75 (672GWA),5079-76 (672RBWA), 5079-67 (663BKWA), 5079-68 (663RWA), 5079-70 (663GWA), 5079-71 (663BWA),*
meet, (where applicable), the provisions of Council Directive 93/42/EEC *(Annex VII),* and pertaining to medical devices.

This Declaration is only applicable to the products bearing the product numbers listed above that have been processed with WI 7.6-2 CE Marked Products and that have records maintained in WI 4.5-3 Device History Records.

The product is classified as Class I according to the European Directive 93/42/EEC.

We hereby appoint ADC LTD., Unit 6 Fareham Enterprise Centre, Newgate Lane, Hants, UK, PO14 1TH, as European Responsible Person as stipulated in Art. 14.2 of the Medical Device Directive 93/42/EEC.

Limitations to this Declaration include:
None

For all of the applicable standards for this device, please reference Section II of Annex D of the Stethoscopes Technical File.

Signed this day 5 Of December 2013

Mike Falco
Quality Manager

The compliance with requirements of *Annex VII* have been approved by TUV Rheinland LGA Products GmbH (EU Identification No. 0197) Am Grauen Stein D-51105 Köln.

This document was created on *12/5/2013*, at American Diagnostic Corporation, 55 Commerce Drive, Hauppauge, NY, 11788, USA.

CE Certificate Number: DD 60037651 0001

Internal Certificate Number: *12052013STETH*

Declaration of Conformity in accordance with ISO/IEC 17050-1.

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