TÜVRheinland®

EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60090676 0001

Report No.: 15062915 001

Manufacturer:

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

Products:

Aspects of manufacture concerned with conformity of products with the metrological requirements of Aneroid Sphygmomanometers and Mercury Sphygmomanometers restricted for Healthcare Use only

Replaces Approval, Registration No.: DD 60035984 0001

Expiry Date: 2018-11-01

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2013-12-10

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TÜV Rheinland LGA Products GmbH Zertifizierungsstelle
TÜVRheinland®
Notified Body
Ren

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.