EC Certificate

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

Registration No.: DD 60112934 0001

Report No.: 15057479 007

Manufacturer:

Wuxi Medical Instrument Factory
No. 43, Xinxi Road
Zhangjing, Xibei Town
Wuxi City
214194 Jiangsu
China

Products:

Alcohol Pads;

Aspects of manufacture concerned with conformity of products with the metrological requirements of Sphygmomanometers

Expiry Date: 2021-08-08

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.

Notified Body

Effective Date: 2016-08-09

Date: 2016-08-09

X. 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.