TÜVRheinland

EC Certificate

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

Registration No.: DD 60113351 0001

Report No.: 15044592 007

Manufacturer:
Wenzhou Jianda Medical
Instrument Co., Ltd.
661 Xiayang St.
Yongxing Wenzhou
325024 Zhejiang
China

Products:
Aspects of manufacture concerned with conformity of products with the metrological requirements of Aneroid Sphygmomanometers

Replaces Approval, Registration No.: DD 60040552 0001

Expiry Date: 2021-10-14

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: 2016-10-28

Date: 2016-10-28

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.