EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4
Full Quality Assurance System
Medical Devices

Registration No.: HD 60139416 0001

Report No.: 17029954 012

Manufacturer:
Shenzhen Jiacom Technology
Co., Ltd.
301, No.596-4 Dahe Village
Guancheng Community
Guanhu Street, Longhua District
518110 Shenzhen
P.R. China

Products:
- Blood Pressure Monitors
- Infrared Thermometers
- Compressor Nebulizers
- Vital Sign Monitors

Replaces Approval, Registration No.: HD 60127617 0001

Expiry Date: 2023-04-29

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 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 II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date: 2020-03-18

Date: 2020-03-18

Notified Body

Dipl.-Ing. I. Munkler

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
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