EC Certificate

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

Registration No.: HD 60123955 0001

Report No.: 17039584 005

Manufacturer:
Shenzhen Viatom Technology
Co., Ltd.
4E, Building 3, Tingwei Industrial Park
No. 6 Liufang Road, Block 67
Xin'an Street, Baoan District
Shenzhen
518101 Guangdong
China

Products:
- Vital Signs Monitors
- Pulse Oximeters
- Blood Pressure Monitors

Replaces Approval, Registration No.: HD 60114302 0001

Expiry Date: 2019-09-01

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: 2018-02-13

Date: 2018-02-13

Notified Body
TÜV Rheinland LGA Products GmbH
TÜVRheinland
III
Sertifizierungsstelle
S. Liu

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.