

**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

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**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.