

# EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: **Shenzhen Viatom Technology Co., Ltd.  
4E, Building 3, Tingwei Industrial Park,  
No.6 Liufang Road, Block 67, Xin'an Street,  
Baoan District, 518101 Shenzhen, P.R.China**

Name and address of Authorized Representative: **Well Kang Limited  
The Black Church, St. Mary's Place, Dublin 7, D07  
P4AX, Ireland  
TEL1: +353 (1) 2542900  
TEL2: +353 (1) 4433560  
FAX: +353 (1) 6864856**

We declare that the product concerned has been designed and manufactured under a quality management system under a quality management system according to Annex IX of directive 93/42/EEC.

the medical device: **Blood Pressure Monitor  
Model:  
BP1, BP1A  
Include accessories  
Cuff: CU-10**

UMDNS of class: **16173  
Class IIa**

Conformity assessment procedure: **MDD 93/42/EEC Annex II excluding (4)**

Certificate No.: **HD 60137356 0001**

Issue date: **2019-07-17**

Expiry date: **2024-05-27**

Notified Body: **TÜV Rheinland LGA Products GmbH  
Tillystraße 2  
90431 Nürnberg  
Deutschland  
CE 0197**

This Declaration of Conformity covers all medical devices as specified in the product list belonging to this declaration and is only valid in connection with a batch specific Certificate of Compliance for all products concerned bearing the CE mark.

Shenzhen, 2020/06/03  
Place, date

Management representative Wang Guannan  
Name and function

