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,Dublin7,D07 P4AX,Ireland** **TEL1: +353（1）2542900** **TEL2: +353（1）4433560** **FAX: +353（1）6864856**

We declare under our sole responsibility 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, BP2, BP2A** **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