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, Shenzhen 518101 Guangdong China**
Name and address of Authorized Representative:	**MedNet EC-REP GmbH** **Borkstrasse 10 , 48163 Muenster,Germany** **Telefon: +49 251 32266-61** **Telefax: +49 251 32266-22**

We declare under our sole responsibility that

the medical device:	**Pulse Oximeter** **Model: Oxiband， PO6， PO4，PO5 ， PO2**
UMDNS	**17148**
of class:	**Class IIa**
	according to annex IX of directive 93/42/EEC
Conformity assessment procedure:	**MDD 93/42/EEC Annex II excluding (4)**
Conformity assessment procedure:	**Directive 93/42/EEC Annex II.3**
Registration No.:	**HD 60137356 0001**
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, 2020/11/04
Place, date

General Manager Zhou Saixin
Name and function