# EC DECLARATION OF CONFORMITY

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| Name and address of the manufacturer: | **Shenzhen Viatom Technology Co., Ltd.**<br>**4E,Building 3, Tingwei Industrial Park,**<br>**No.6 Liufang Road, Block 67, Xin'an Street,**<br>**Baoan District, Shenzhen 518101 Guangdong China** |
| Name and address of Authorized Representative: | **MedNet EC-REP GmbH**<br>**Borkstrasse 10 , 48163 Muenster,Germany**<br>**Telefon: +49 251 32266-61**<br>**Telefax: +49 251 32266-22** |

We declare under our sole responsibility that

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| the medical device: | **Pulse Oximeter**<br>**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)**

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