

**Original Declaration of Conformity**

**EU DECLARATION OF CONFORMITY**

**Apparatus/Equipment**

Product: Electronic Upper Arm Blood pressure monitor MT-30  
 Model: KD-5812(MT-30)  
 SKU No.: N/A  
 Batch or Serial Number: N/A

**Manufacturer or his authorized representative:**

Name: Andon Health Co., Ltd.  
 Address: No. 3 JinPing Street, YaAn Road,  
 Nankai District, Tianjin 300190  
 Country: China

**Manufacturers authorized EU Representative**

Name: iHealthLabs Europe SAS  
 Address: 36 Rue de Ponthieu, 75008, Paris  
 Country: France

**This declaration of conformity is issued under the sole responsibility of the manufacturer.**

Object of the declaration: Listed above

The object of the declaration described above is in conformity with the following relevant Union harmonization directives and/or legislation(s):

Radio Equipment Directive (RED)	2014/53/EU	<input checked="" type="checkbox"/>
Low Voltage Directive (LVD)	2014/35/EU	<input type="checkbox"/>
Electromagnetic Compatibility Directive (EMCD)	2014/30/EU	<input type="checkbox"/>
Restriction of the use of certain hazardous substances (RoHS)	2011/65/EU	<input type="checkbox"/>
Ecodesign Requirements for energy-related products (ErP)	2009/125/EC	<input type="checkbox"/>
Machinery (MD)	2006/42/EC	<input type="checkbox"/>
Toys safety	2009/48/EC	<input type="checkbox"/>
Personal protective equipment (PPE)	Regulation (EU) 2016/425	<input type="checkbox"/>
Construction products (CPD/CPR)	Regulation (EU) No 305/2011	<input type="checkbox"/>
Cosmetic products	Regulation (EC) No 1223/2009	<input type="checkbox"/>
Medical devices(MDD)	Directive 93/42/EEC	<input type="checkbox"/>
		<input type="checkbox"/>

References to the relevant harmonized standards used, including the date of the standard, or references to the other technical specifications, including the date of the specification, in relation to which conformity is declared:

Harmonized Standard	Test Report No.	Test lab
EN 62368-1:2014+A11:2017	S202206015801S01	Fanguang Inspection & Testing Co.,Ltd.
ETSI EN 301489-1 V2.2.3(2019-11) ETSI EN 301489-17 V3.2.4(2020-09) EN 55032:2015+A11:2020 EN 55035:2017+A11:2020 EN IEC 61000-3-2:2019 EN 61000-3-3:2013+A1:2019	S202206015801E01	Fanguang Inspection & Testing Co.,Ltd.
ETSI EN 300 328 V2.2.2 (2019-07)	S202206015801E02	Fanguang Inspection & Testing Co.,Ltd.

**Notified Body (Optional) :**

Name of notified body: MICOM Labs Inc  
 Reference Number of the certificate: 2280  
 4 digit notified body number:

Signed for and on behalf of Peroxfarma S.A.  
 Place and date of issue: Tianjin,China, October 12, 2022

Signature: Sun Guimei

Name, function: SunGuimei Certificate Engineer



Color Image of product:

