

# SCHEDA TECNICA

Codice	SFI230
Modello:	U80EH
Contenuto confezione:	Sfigmomanometro digitale da braccio; manuale istruzioni





# Caratteristiche Tecniche

Descrizione	Sfigmomanometro digitale da braccio per la misurazione della pressione sanguigna		
Display	LCD display		
Principio di misurazione	Metodo Oscillometrico		
Posizione di misurazione	Parte superiore del braccio		
Intervallo di misurazione	Pressione:	0~299 mmHg	
	Pulsazioni:	40~199 pulsazioni/minuto	
Precisione	Pressione:	±3mmHg	
	Pulsazioni:	±5% of reading	
Indicazioni LCD Pressione: 3 digits display of mmHg		3 digits display of mmHg	



	Pulsazioni:	3 digits display	
	Simboli:	Memoria/ Battito cardiaco/Batteria bassa	
Memoria	2x90 misurazioni memorizzabili		
Alimentazione	4 Batterie Alcaline AA DC.6V o adattatore AC		
Spegnimento automatico	in 3 minuti		
Peso uniità	Approssimativamente 219g (batterie non comprese)		

Dimensione	132mm×100mm×45mm		
Usura apparecchio	10,000 misurazioni in condizioni normali		
Vita batteria	Circa 300 misurazioni in condizioni normali		
Accessori	Bracciale, manuale di istruzioni		
Condizioni di utilizzo	Temperatura:	5°C~40°C	
	Umidità:	15%~93%RH	
	Pressione atmosferica:	86kPa~106kPa	
Condizioni di stoccaggio:	Temperatura:		-20°C~55°C
	Umidità:		10%~93%RH;
Evitare urti, esposizione a raggi solari e a pioggia durante il			
trasporto	Pressione atmosferica:		86kPa~106kPa



Si consiglia di leggere attentamente il manuale di istruzioni

# PVS SpA – Cassina de Pecchi IT - Rev. 03/03/2021







td.

# **EC** Certificate

Full Quality Assurance System Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III) **No. G1 078672 0014 Rev. 00** 

**Manufacturer:** 

Shenzhen Urion Technology Co., L Floor 4-6th of Building D

Jiale Science & Technology Industrial Zone No.3, ChuangWei Road Heshuikou Community, MaTian Street GuangMing New District 518106 Shenzhen PEOPLE'S REPUBLIC OF CHINA

# Product Category(ies): Digital Blood Pressure Monitors, Infrared Thermometer, Compressor Nebulizers

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

GZ1917901

Valid from: Valid until: 2020-02-21 2022-02-07

Date,

2020-02-21

Christoph Dicks Head of Certification/Notified Body

A4 / 07.17





# **EC Certificate**

Full Quality Assurance System Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (Devices in Class IIa, IIb or III) No. G1 078672 0014 Rev. 00

Facility(ies):

Shenzhen Urion Technology Co., Ltd.

Floor 4-6th of Building D, Jiale Science & Technology Industrial Zone, No.3, ChuangWei Road, Heshuikou Community, MaTian Street, GuangMing New District, 518106 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

(4



Page 2 of 2 TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany



# Upper Arm Electronic **Blood Pressure Monitor**

Model:U80EH



# Instruction Manual

# **Table of Contents**

Introduction ·····	03
Safety information ·····	04
Product structure	08
-Each part Name	08
Battery Installation	09
Setting mode	11
-How to set	11
Proper use of the unit	13
-Pre-measurement ·····	13
-Common factors of wrong Measurement	13
-Fitting the cuff	14
-Measuring procedure	15
-Discontinuing a Measurement	15
-Memory-recall of measurements	15
Reading memory record	16
Memory -clear of measurements	16
About blood pressure	16
Exceptional situations	18
Care and maintenance	19
Specifications	20
Warranty information	21
EMC Declaration	22
02/25	

# Introduction

▲ Your new digital blood pressure monitor uses the oscillometric method of blood pressure measurement. This means the monitor detects your blood's movement through your brachial artery and converts the movements into a digital Reading. An oscillometric monitor does not need a stethoscope, so the monitor is simple to

▲ Intelligent inflation will reduce the uncomfortable feeling by incorrect inflation, and shorten the measurement time, prolong the cuff's usage lifetime

▲ 2x90 sets memory function, each measurement result will be displayed on the screen, and automatically stored. This unit has blood classification index, could easy to check your blood pressure

Please read the manual carefully before you use the unit, and keep the manual well after using.

#### CONTRAINDICATION

This product can't be used in patients who is with severe heart insufficiency to avoid suffocation and death. This product is not suitable for infants and children.

# INTENDED USE

The automatic blood pressure monitor intended to measurement the systolic pressure, diastolic pressure and pulse rate through upper arm.They are expect used into the home and hospital, intended for over than 12 years old adult using.

# Safety Information

To assure the correct use of the product, basic safety measures should always be followed including the warning and the caution listed in the instruction manual

# Symbol descriptior

The following symbols may appear in this manual, on the label, on the device, or on it's accessories. Some of the symbols represent standards and compliances associated with the device and its use.

MARNING: This alert identifies hazards that may cause serious personal injury or death

A CAUTION: This alert identifies hazards that may cause

minor personal injury, product damage, or property damage

- Type BF applied part
- Manufacturer **SN** Specifies serial number
- EC REP Authorized Representative in the European Community
- $\mathbf{C} \in \mathbf{C}$  CE Mark: conforms to essential requirements of the Medical Device Directive 93/42/EEC.
- DISPOSAL: Do not dispose this product as unsorted municipal waste. Collection of such waste separately
- for special treatment is necessary. === Direct current
- Follow instructions for use
- ▲ CAUTION: Consult accompanying documents

04/25

# Safety Information

▲ Those who have arrhythmia, diabetes, blood circulation or apoplexy problem, please use under the physician's instruction. ▲ Contact your physician for specific information about your blood pressure.Self diagnosis and treatment which use measured results

may be dangerous. Follow the instructions of your physician or licensed healthcare provider. A Please place on a high place where children can't be touched.

▲ No modification of this equipment is allowed.

A Do not modify this equipment without authorization of the manufacturer.

▲ If this equipment is modified, appropriate inspection and testing must be conducted to ensure continued safe use of equipment. ▲ The cuff hose around neck may cause the suffocation

A The swallowing of samll park like packing bag, battery, battery cover and on may cause the suffocation.

 $\triangle$  Please don't use a dilution agent, alcohol or petrol to clean the unit.Please don't hit heavily or fall down the product from a high place.Use the right cuff, otherwise it can not work.

 $\triangle$  Never leave any low battery in the battery compartment since they may leak and cause damage to the unit.

 $\triangle$  Please take off the battery if you won't use in 3 months.  $\underline{\Lambda}$  Replace the new batteries if the unit display a low battery symbol.

 $\underline{\wedge}$  Do not mix the old and new batteries.

▲ Do not use a cellular phone near the unit.It may result in operational failure.

▲ Please avoid using in high radiant area in order to make your measuring data correctly

05/25

# Safety Information

 $\underline{\Lambda}$  Consecutive blood pressure measurements should be repeated after 1 minute pause or after the arm has been held up in order to allow the accumulated blood to flow away.

 $\Lambda$  If the arm circumference size is beyond the measuring range of CUFF, it can't be measured and used, then it will cause the blood flowing unsmooth and wrong measurement data.

▲ Don't kink the connection tube during use, otherwise the cuff pressure may continuously increase which can prevent blood flow and result in harmful injury to the PATIENT.

 $\triangle$  Too frequent measurements can cause injury to the PATIENT due to blood flow interference.

 $\triangle$  Don't apply CUFF over a wound, it can cause further injury to the PATIENT.

The device is not suitable for use on neonatal patients, pregnant  $\mathbf{\Lambda}$ women, patients with implanted, electronical devices, patients with pre-eclampsia, premature ventricular beats, atrial fibrillation, peripheral, arterial disease and patients undergoing intravascular therapy or arterio-venous shunt or people who received a mastectomy. Please consult your doctor prior to using the unit if you suffer from ilnesses.

 $\triangle$  When using this device, please pay attention to the following situation which may interrupt blood flow and influence blood circulation of the patient, thus cause harmful injury to the patient: connection tubing kinking too frequent :the application of the cuff and its pressurization on any arm where intravascular access or therapy, or an arterio-venous (A-V) shunt, is present; inflating the cuff on the side of a mastectomy.

 $\wedge$  Do not inflate the cuff on the same limb which other monitoring ME equipment is applied around simultaneously, because this could cause temporary loss of function of those.

 $\underline{\Lambda}$  Please check that operation of the device does not result in prolonged impairment of patient blood circulation.

# Safety Information

 $\Lambda$  Do not use the equipment where flammable gas(such as anesthetic gas, oxygen or hydrogen) or flammable liquid (such as alcohol)are present

▲ Do not touch the output of AC adapter and the patient simultaneously

 $\underline{\Lambda}$  Do not touch the live end of battery and the patient simultaneously when change the batteries

# A WARNING:

Do not dispose of electrical appliances as unsorted municipal waste, use separate collection facilities.Contact you local government for information regarding the collection systems available. If electrical appliances are disposed of in landfills or dumps, hazardous substances can leak into the groundwater and get into the food chain, damaging your health and well-being

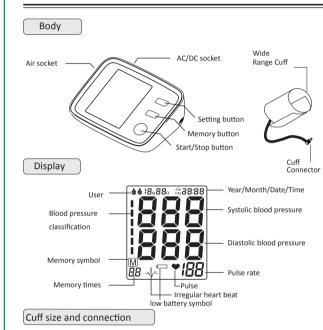
#### Classification

- 1. Internally powered equipment;
- 2. Type BF applied part; .
  - 3. Protection against ingress of water or Particulate matter: IP21; 4. Not category AP /APG equipment;
  - 5. Mode of operation: Continuous operation.

 $\mathbf{A}$  The user must check that the equipment functions safely and see that it is in proper working condition before being used

07/25

# Product structure



The accessories cuff is M size, for upper-arm circumference 22-32cm use.The cuff is treated as the applied part.

Insert the connector with cuff tube into the hole which is on the left

08/25

side of the device as picture. (Only provided cuff can be used, can not change to any other branded cuff.)

# **Battery installation**

#### Battery installation

the proper direction.

showed.

Remove the battery cover from the battery compartment, insert the battery. a)Remove the battery cover as picture

b)Insert 4 AA powerful batteries into the compartment and ensure each battery is in

#### Low battery and replacement

When power on, the low battery symbol 📼 will display once the unit start to work, and you must re lace with new b unit can' t work.

#### Battery type and replacement

Please use 4pcs AA identical 1. 5V alkaline batteries. Do not use the batteries beyond their expiry date. Please remove the batteries if you do not need to use for long time.

#### WARNING

Dispose of the battery in accordance with all federal. state and local laws. To avoid fire and explosion hazard, do not burn or incinerate the battery.

09/25

# enter into the time setting mode.

# <u>) (0 10 20 20</u> Month setting

# 4.Time setting

Continue to above step, the screen will display xxMxxD and xx:xx, and keep flashing on the digits of hour, the digit will increase 1 when press button MEM each time, you could choose from 0 to 23. Press button SET when you confirm the hour, then the digits of minute start to flash, same as the hour setting, each time you press button MEM the digits will keep changing from 00 to 59. Press button SET when you confirm the minute, then the total setting mode is completed.



# **Battery installation**

# Adapter usage(option)

1.When optional AC adapter should comply with the requirement of IEC 60601-1:2005. Furthermore all configurations shall comply with the requirements for medical electrical systems(see IEC 60601-1-1 or clause 16 of the 3Ed.of IEC 60601-1, respective-ly). Anybody connecting additional equipment to medical electrical equipment configures a medical system and is therefore responsible that the system complies with the requirements for medical electrical systems. Attention is drawn to the fact that local laws take priority over the above mentioned requirements.If in doubt, consult your local representative or the technical service department.

2. This device is double insulated and protected against short circuit and overload by a primary thermal fuse. Make sure to take the batteries out of the compartment before using the mains part. Equipment class 2. 3.When using AC power, to avoid possible damage to the monitor, use only the exclusive AC adapter that can be purchased from authorized dealers. 4.Insert the adapter plug into the hole on the backside of the unit as picture.

5.Insert the other side of the adapter into the outlet with 100-240V. 6.To remove the AC adapter, disconnect the adapter plug from the outlet first and then disconnect the cord from the unit's socket.

Adapter technical features: Output voltage:6V±5%

Output current:At least 600 mA

Output plug polarity: <+> inner

and restart the measurement.

å

USER 1

Setting mode

Note:

HOW TO SET

1.user setting

2.Year setting

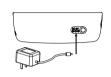
Setting mode

3.Month and date setting

mode

External diameter: 5. 5mm 0.1 mm

Internal diameter: 2.1 mm 0.1 mm



10/25

When use AC adapter, the power of battery won't be consumed. •When suddenly stop during measurement(like the plug off from the outlet by carelessness), it must be reinserted the plug into the unit,

Press button SET when power off, the screen will display norman , press button MEM, it will be changed between and and a, press button SET when you confirm the user, then it will enter into the year setting



Continue to above step, the screen will display and flash 20XX, the last digit of the year will increase 1 when press button MEM each time, you could choose from 2020 to 2099.Press button SET when you confirm the year, then it will enter into the month and date setting mode.

Year setting

11/25

#### Continue to above step, the screen will display xxMxxD and xxxx, and keep flashing on month, the digit will increase 1 when press button MEM each time, you could choose from 1to 12.Press button SET when you confirm the month, then it will set the date. Same as the month setting, each time you press button MEM, the digit will keep changing from 01 to 31.Press button SET when you confirm the date, then it will

606 2020





# Proper use of the unit

# Measurement

Pre-measurement

·Relax for about five to ten minutes prior to the measurement Avoid eating, drinking alcohol, smoking, exercising and bathing for 30 minutes before taking a measurement.

- All these factors will influence the measurement result.
- ·Remove any garment that fits closely to your upper arm.
- ·Always measure on the same arm(normally left)

•Take measurement regularly at the same time of every day, as blood pressure changes even during the day.

# Common factors of wrong measurement

 $\cdot All$  efforts by the patient to support their arm can increase blood pressure

·Make sure you are in a comfortable, relax position and do not activate any of the muscles in the measurement arm during measurement. Use a cushion for support if necessary.

·If the arm artery lies lower or higher than the heart, a false reading will be obtained.

#### Note

•Only use clinically approved cuffs!

·A loose cuff or a exposed bladder causes false reading.

·With repeated measurements.blood accumulates in the arm which can lead to false reading.

Consecutive blood pressure measurements should be repeated after 1 minute pause or after the arm has been held up in order to allow the accumulated blood to flow away.

13/25

# Proper use of the unit

#### **Fitting the CUFF**

1). Put the cuff on a table flatly with the velcro side down.Pass the end of the cuff through the metal loop so that a circle is formed The velcro closer will now be facing outwards (ignore this step if the cuff has already been prepared).

2). Push the cuff over the left upper arm so that the tube points in the direction of the lower arm. 3). Wrap the cuff on the arm as illustrated Make

certain that the lower edge of the cuff lies approximately 2 to 3 cm above the elbow and the rubber tube leaves the cuff on the inner side of the arm.

4). Tighten the free end of the cuff and close the cuff by affixing the velcro.

5). The cuff should be snug on your upper arm so That you can fit 2 fingers between the cuff and your upper arm. Any piece of clothing restricts the arm which must be taken off.

6). Secure the cuff with the velcro closer in such a way that it lies comfortably and not too tight.Lay your arm on a table(palm upwards)so that the cuff is at the same height as the heart.Do not bend the tube

Note

If it is not possible to fit the cuff to your left arm, it can also be placed on the right. However, all measurements should be made using the same arm

14/25

# Proper use of the unit

# Measuring procedure:

After the cuff has been appropriately positioned, the measurement can begin:

1). Press the START/STOP button, all symbols appear on the display, then the pump begins to inflate the cuff, the rising pressure in the cuff is shown on the display. 2). After the suitable pressure has been reached, the pump stops and the pressure gradually falls. The cuff pressure is displayed. In case that the inflation is not sufficient, the device automatically re-inflates to a higher pressure.

3). When the device detects the signal, the heart symbol 🎔 on the display starts to flash. 4). When the measurement has been completed, the systolic, diastolic and pulse rate will appear on the

display. 5). The measurement readings remain on the display until you switch off the device. If no button is pressed for a period of 3 minutes, the device switches off itself in order to save the power.

Note: The symbol  $\sqrt[n]{}$  will be displayed along with the reading if the irregular (B)heartbeat is detected during the measurement.

#### Discontinuing a measurement

If it is necessary to interrupt a blood pressure measurement for any reason(eg.the patient feels unwell)the START/STOP button can be pressed at any time. The device immediately decrease the cuff pressure automatically.

# Memory-recall of measurements

This blood pressure monitor automatically stores 2×90 sets measurements value, the oldest record will be replaced by the latest measurement value when more than 90 sets each user. 15/25

A Row 5st - At-



150



# About blood pressure

#### Read memory record

Press the button MEM when power off the latest 3 times average value will be shown, press the button MEM again, the last measurement value will be shown, as well as subsequent measurements can be display one after the other by pressing the button MEM each time



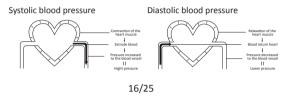
#### Memory -clear of measurements

If you are sure that you want to permanently remove all stored memories.Press the button SET for 7 times until CL appears when power off, press the START/STOP button, CL will flash for 3 times to clear all the memories. After this press button MEM, M and "no" will be shown on the display which mean that no memory in store.

# About blood pressure

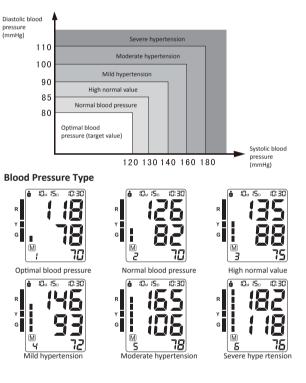
Blood pressure is the pressure exerted the arteries. The systolic blood pressure value represents the blood pressure produced by contraction of the heart muscle.

The diastolic blood pressure value represents the blood pressure produced by relaxation of the heart muscle.



#### About blood pressure

According to the blood pressure classification by the WHO/ISH. SYS lower than 100mmHg is considered as hypotension.



17/25

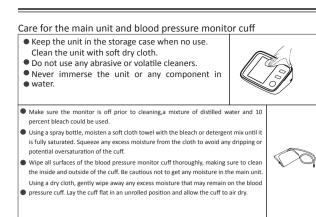
# **Exceptional Situation**

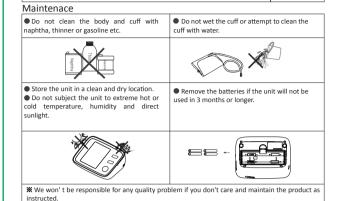
Error indicators The following symbol will appear on the display when measuring

Symbol		Cause	Correction			
Symbol			Correction			
Weak signal or F -   pressure change			Wrap the cuff properly.			
L '	sudd		Remeasure with correct	way.		
E-2 External strong		0	When near cell phone or other high radiant device the measurement will be failed.			
	aisea	bunce	Keep quite and no chattin	Keep quite and no chatting when measure.		
		pears error	Wrap the cuff properly.			
8-3		ig the	Make sure that the air plug is properly inserted in the unit.			
C-J	process of inflating		Remeasure.			
8-5	Abno press	rmal blood ure	Repeat the measurement after relax for 30 mins, if get unusual readings for 3 times, please contact your doctor.			
	low	oattery	Replace all the worn batteries with new ones.			
rouble re	mov	al				
Problem			Check	Cause and solutions		
No power		Check the battery power		Replace new one		
No power		Check the polarity position		Installation for proper placement of the batteries polarities		
No inflation		Whether the plug insert		Insert into the air socket tightly		
		Whether the plug broken or leak		Change a new cuff		
Err and stop working		Whether move the arm when inflate		Keep the body peaceful		
		Check if chatting when measured		Keep quite when measure		
Cuff leak		Whether the cuff wrap too loose		Wrap the cuff tightly		
		Whether the cuff broken		Change a new cuff		
		whether the	cull broken	change a new cun		

18/25

# Care and maintenance





19/25

#### Specification

#### Automatic upper arm blood pressure monito Description Display LCD digital display Measuring principle Oscillometric method Measuring localization Upper arm Measuremen Pressure 0~299mmHg range 40~199 pulses/min Pulse Pressure ±3mmHg Accuracy Pulse ±5% of reading Pressure 3 digits display of mmHg LCD Pulse 3 digits display indication symbol Memory/Heartbeat/Low battery 2x90 sets memory of measurement values Memory function 4pcs AA alkaline battery DC.6V or AC adapter Power source Automatic power off in 3 minutes Approx.219g(batteries not included) Main unit weight Main unit size 132mm×100mm×45mm Main unit lifetime 10,000 times under normal use Battery life Could be used for 300 times for normal condition Accessories Cuff, instruction manual Temperature 5°C~40°C Operating 15%~93%RH Humidity environment Air pressure 86kPa~106kPa Air pressure:86kPa~106kPa; Storage Temperature:-20°C~55°C; Humidity:10%~93%RH; environmen avoid crash, sun burn or rain during transporta

20/25

# Warranty information

#### Statement

- The intended use: the unit is intended to be used by adults at home or medical center to measure blood pressure and pulse rate from the wrist.
- The unit satisfies the requirements of EN ISO 81060-1 Part 1 Noninvasive sphygmomanometers, EN 1060-3:1997+ A2:2009 Non-invasive sphygmoma-
- nometers. IEC80601-2-30 Part 2 Non-invasive sphygmomanometers. Blood pressure measurements determined with this device are equivalent to
- those obtained by a trained observer using ■ The cuff stethoscope auscultatory method, within the limits prescribed by
- the American National Standard, manual, electronic, or automated sphygmomanometers.
- The risk of patient and user can be lowered to acceptable level.

### Warranty Information

- The unit is guaranteed to be free of defects in workmanship and materials under normal use for a period of Two Years from the date listed on the purchase record.
- For repair under this warranty. Our authorized service agent must be advised of the fault with the period of the warranty. This warranty covers parts and labor only under normal operations. Any defect resulting from natural causes, eg. flood, hurricane etc, is not within this guarantee. This guaranty does not cover damage incurred By use of the unit not in accordance with the instructions, accidental damage, or being tampered with or serviced by unauthorized service agents.
- Monitor subjected to misuse, abuse, and neglect of these manual content.non-instructional purposes:unauthorized repair or modifications will be excluded from this warranty.

21/25

- The device requires no calibration.
- ▲ The device is not repairable and contains no user serviceable parts.

# **EMC Declaration**

IEC 60601-1-2: 2014 ME EQUIPMENT and ME SYSTEMS identification, marking and documents for Class B product

Instructions for use The ME EQUIPMENT or ME SYSTEM is suitable for home healthcare environ-

ments and so on. Warning: Don't near active HF surgical equipment and the RF shielded room of an ME system for magnetic resonance imaging, where the intensity of EM

disturbances is high. Warning: Use of this equipment adjacent to or stacked with other equipment

should be avoided because it could result in improper operation. If such use is necessary, this equipment and the other equipment should be observed to verify that they are operating normally.

Warning: Use of accessories, transducers and cables other than those specified or provided by the manufacturer of this equipment could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation."

Warning: Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the blood pressure monitor, including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.

If any: A list of all cables and maximum lengths of cables (if applicable), transducers and other ACCESSORIES that are replaceable by the RESPONSIBLE ORGANIZATION and that are likely to affect compliance of the ME EQUIPMENT or ME SYSTEM with the requirements of Clause 7 (EMISSIONS) and Clause 8 (IMMUNITY). ACCESSORIES may be specified either generically (e.g. shielded cable, load impedance) or specifically (e.g. by MANUFACTURER and EQUIPMENT OR TYPE REFERENCE).

If any: The performance of the ME EQUIPMENT or ME SYSTEM that was determined to be ESSENTIAL PERFORMANCE and a description of what the OPERATOR can expect if the ESSENTIAL PERFORMANCE is lost or degraded due to EM DISTURBANCES (the defined term "ESSENTIAL PERFORMANCE" need not be used).

22/25

# **EMC** Declaration

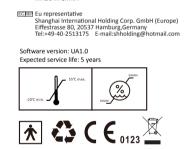
Technical description 1.All necessary instructions for maintaining BASIC SAFETY and ESSENTIAL

PERFORMANCE with regard to electromagnetic disturbances for the excepted service life

2. Guidance and manufacturer's declaration -electromagnetic emissions and Immunity

Tabe 1

Guidance and manufacturer's declaration - electromagnetic emissions		
Emissions test	Compliance	
RF emissions CISPR 11	Group 1	
RF emissions CISPR 11	Class B	
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Compliance	



# **EMC** Declaration

Guidance and mar	nufacturer's declaration - electroma	gnetic Immunity	
Immunity Test	IEC 60601-1-2 Test level	Compliance level	
Electrostatic discharge (ESD) IEC 61000-4-2	±8 kV contact ±2 kV, ±4 kV, ±8 kV, ±15 kV air	±8 kV contactv ±2 kV, ±4 kV, ±8 kV, ±15 kV air	
Electrical fast transient/burst IEC 61000-4-4	Power supply lines: ±2 kV 100 kHz repetition frequency	Power supply lines: ±2 kV 100 kHz repetition frequency	
Surge IEC 61000-4-5	line(s) to line(s): ±0.5kV ±1 kV	line(s) to line(s): ±0.5kV ±1 kV	
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	0% 0.5 cycle At 0°, 45 °, 90 °, 135 °, 180 °, 225 °, 270 ° and 315 ° 0% 1 cycle And 70% 25/30 cycles Single phase: at 0% 300 cycle	0% 0.5 cycle At 0º, 45 º, 90 º, 135 º, 180 º, 225 º, 270 º and 315 º 0% 1 cycle And 70% 25/30 cycles Single phase: at 0% 300 cycle	
Power frequency magnetic field IEC 61000-4-8	30 A/m 50Hz/60Hz	30 A/m 50Hz/60Hz	
Conduced RF IEC61000-4-6	150KHz to 80MHz: 3Vrms 6Vrms (in ISM and amateur radio bands) 80% Am at 1kHz	150KHz to 80MHz: 3Vrms 6Vrms (in ISM and amateur radio bands) 80% Am at 1kHz	
Radiated RF IEC61000-4-3	10 V/m 80 MHz – 2,7 GHz 80 % AM at 1 kHz	10 V/m 80 MHz – 2,7 GHz 80 % AM at 1 kHz	

23/25

NOTE UT is the a.c. mians voltage prior to application of the test level.

# 24/25

# EM