telli health

Arm Blood Pressure Monitor

Model: TelliBP01

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- converts the movements into a digital reading. An oscillometric
- ▲ Intelligent inflation will reduce the uncomfortable feeling by incorrect inflation, and shorten the measurement time, prolong
- ▲ 2x90 sets memory function,each measurement result will be displayed on the screen, and automatically stored. This unit has
- ▲ Please read the manual carefully before you use the unit, and keep the manual well after using.

This product can't be used in patients who is with severe heart insufficiency to avoid suffocation and death

This automatic blood pressure monitor intends to measure the systolic pressure, diastolic pressure and pulse rate through upper arm. It's expected to be used at home or in the hospital, intended for people over 12 years old.

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:

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

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

- ⚠ 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 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

Safety Information

⚠ Those who have arrhythmia, diabetes, blood circulation or

apoplexy problem, please use under the physician's

⚠ 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

⚠ If this equipment is modified, appropriate inspection and

The cuff hose around neck may cause the suffocation.

battery cover and so on may cause the suffocat

of device with rechargeable lithium battery).

⚠ The swallowing of small part like packaging bag, battery,

⚠ 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

⚠Do not replace or remove the battery from device (in the case

⚠ Do not use a cellular phone near the unit. It may result in

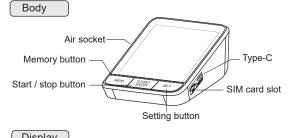
APlease avoid using in high radiant area in order to make

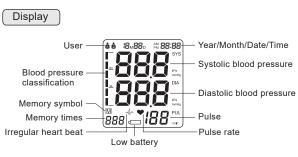
testing must be conducted to ensure continued safe use of

A Please place on a high place where children can't be

A No modification of this equipment is allowed.

Product structure





Cuff size and connection

Battery installation

SIM card installation

The accessories cuff is universal size, for upper-arm circumference 22-42 cm use. The cuff is treated as the applied part.

Insert the connector with cuff tube into device as picture. (Only provided cuff can be used, can not change to any other branded cuff.)

Inserted the SIM card into the card slot on the right

1.Remove the battery cover from the battery compartment,

Positive (+) and Negative (-) are displayed on the device

side of device before measurement. The card

should be in accordance with direction "

and chip side face down. When you need

to take out the card, just press the card

following the arrow for proper removal

18650 is a standard lithium-ion battery

diameter of 18mm, 65 represents a

length of 65mm, and 0 represents a

model, where 18 represents a

Low battery and chargea

2. Insert 1pc 3.7V 18650 lithium battery, and

ensure the battery is in the proper direction.

inward and then it will pop out.

Battery installation

3. Close the battery cover.

Battery size

cylindrical battery.

<●

Setting mode

Adapter technical features:

Output voltage: Type-C 5V±5% Max.output current: At least 600 mA

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, respectively). Anybody

connecting additional equipment to medical electrical equipment

configures a medical system and is therefore responsible that the

systems. Attention is drawn to the fact that local laws take priority

over the above mentioned requirements. If in doubt, consult your

2. When using AC power, to avoid possible damage to the monitor,

authorized dealers. Other adapters may vary in output voltage and

use only the exclusive AC adapter that can be purchased from

3. Insert the adapter plug into the hole on the backside of the

4. Insert the other side of the adapter into the outlet

5. To remove the AC adapter, disconnect the adapter

plug from the outlet first and then disconnect the cord

system complies with the requirements for medical electrical

local representative or the technical service department.

Note:

•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, and restart the measurement.

How to set

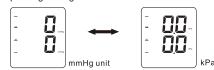
polarities.

unit as picture.

1.Start to set, Unit setting:

Press the SET button when power off, 0 or 0.0 will be displayed, then the setting begin.

Continute to above step, the unit will be changed when press the MEM button each time. Press the SET button to confirm the unit, then it will enter into the Speaking setting mode



2. Speaking setting

Continute to above step, the screen will display SP OF or SP ON, press the MEM button and it will change between SP OF and SP ON, press the SET button to confirm the option. Following this, the device will enter into the User setting mode.

Remark: SP OF: close the saking function.

SP ON: open the speaking function.

Setting mode

5. Year setting

Setting mode

Ūn.

6. Month and date setting

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 1 to 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 enter into the time setting mode.

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

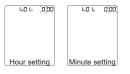
could choose from 2022 to 2099. Press button SET when you confirm

the year, then it will enter into the month and date setting mode.



7. 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.



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 monitor does not need a stethoscope, so the monitor is simple to
- the cuff's usage lifetime
- blood classification index, could easy to check your classification index, could easy to check your

CONTRAINDICATION

This product is not suitable for infants and children.

INTENDED USE

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Safety Information

your measuring data correctly.

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

M 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: Intermittent operation:
- ⚠ The user must check that the equipment functions safely and see that it is in proper working condition before being used.

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Battery installation

the Type-C tip to the USB socket of device.

Noted: the lithium battery could be used for about 310 times when charge is full Charge time is 4-5 hours During charging when power off, the LCD will shows the blood pressure bars on the left side from one bar to six bars in a continuous loop. And unmoved 6 bars along with the battery symbol will be displayed once battery is full' as picture shown

When power on, the low battery symbol will be displayed once the

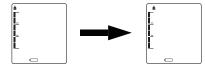
device starts. Plug in the device using the included USB cable to charge.

Take the USB cable and confect it to the charger (charge connector) or

to your USB port on your computer or power bank. Connect the end of

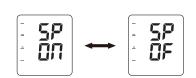


If the device under setting mode and battery charging in the same time, LCD can show the same as above, e.g. user setting



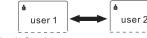
Dispose of the battery in accordance with all federal, state and local laws. To avoid fire and explosion hazard, do not burn or incinerate the battery.

Setting mode



3. User setting

Continue to above step, the screen will display $\hat{\mathbf{n}}$ or $\hat{\mathbf{n}}$, 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 Clock setting mode



4. Alarm clock Setting

Continue to above step, the screen will display "1 OF" and the HOUR digits flash, press the MEM button to change the HOUR digits. Press the SET button to confirm.

Next the MINUTE digits will flash, press the MEM button to change the digits, and press SET button to confirm Next OF or On will flash, press the MEM button to switch between On and

OF. Press the SET to confirm. Once Alarm 1 is set, you can follow the same steps above to set Alarm 2 and Alarm 3.

Remark: The alarms are turned off by default. This device allows you to set three alarm times at which an alarm signal will then be triggered. This can be a useful aid, for example, as a reminder to take your measurements



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

- 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.

- 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.

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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

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



About blood pressure

Signal Strength Indicator

H means signal is strong, L means signal is weak, These marks will appear after finishing measurement and start to upload data

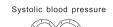




About blood pressure

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



Care and maintenance

Care for the main unit and blood pressure monitor cuff

Clean the unit with soft dry cloth

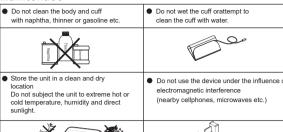
Never immerse the unit or any

Make sure the monitor is off prior to cleaning, a mixture of distilled water and 10 percent bleach could be used.
 Using a spray bottle, moisten a soft cloth towel with the bleach or detergent mix until it is fully saturated. Squeeze any excess moisture from the cloth to avoid any dripping or potential oversaturation of the

cuff.
Wipe all surfaces of the blood pressure monitor cuff thoroughly, making sure to clean the inside and outside of the cuff. Be cautious not to get any moisture in the main unit.
Using a dry cloth, gently wipe away any excess moisture that may remain on the blood pressure cuff. Lay the cuff flat in an unrolled

position and allow the cuff to air dry.

Maintenace



product as instructed

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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 environments and so on.

Warning: Don't near active HF surgical equipment and the RF shielded room of an ME 329mm.n 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 ne cessary, 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 mined 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)

Modulatio Modul Dist IMMUN Freque IEC61000-(W) (m) TEST (Test (V/m) 385 ns for 380 - 390 TETRA 400 27 nodulatior **ENCLOSU** 18 Hz IMMUNITY ± 5 kHz 450 430 -470 kHz sine wireless LTE Band 13 communica 745 704 - 78nodulation tions 780 GSM 800/90

iDEN 820,

CDMA 850,

GSM 1800

CDMA 1900

GSM 1900:

LTE Band 1, 3

WLAN.

802.11 b/g/n

RFID 2450

WLAN 802.11

, 25; UMTS

odulation

Pulse

odulatio

217 Hz

odulation

28

0.3

28

Proper use of the unit

Measuring Procedure:

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

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

Note: The symbol - ✓ will be displyed along with the reading if the irregular heartbeat

Remark: Once finish the measurement, the device it will upload the dada to the background server automatically with showing one rotating signal bar in the top right corner of LCD (as Pic 1).

If Err is displayed (as Pic 2), the upload failed. Upload failure displays Er5 (as Pic 3), indicating that no SIM card is detected. LCD displays nd (as Pic 4) after successful uploading, and automatically shuts down after 5 seconds











150

145

78.





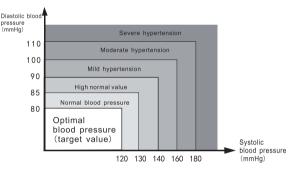
Discontinuing a measurement

natient feels unwell) the START/STOP button can be pressed at any time. The device immediately decrease the cuff pressure automatically

About blood pressure

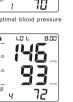
■According to the blood pressure classification by the WHO/ISH.

■SYS lower than 100mmHg (13.3kPa) is considered as hypotension.



■ Blood pressure type







125.

82





88

Specification









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Description	Automatic upper arm blood pressure monitor		
Display	LCD digital display		
Measuring principle	Oscillometric method		
Measuring localization	Upper arm		
Measurement range	Pressure	0~299mmHg	
weasurement range	Pulse	40-199 pulses/min	
Accuracy	Pressure	± 3mmHg	
	Pulse	± 5% of reading	
LCD	Pressure	3 digits display of mmHg	
indication	Pulse	3 digits display	
	symbol	Memory/Heartbeat/Low battery3	
Memory function	2x90 sets memor	2x90 sets memory of measurement values	
Power source	18650 lithium battery 2000mAh/Type-C, working voltage 5V		
Automatic power off	in 3 minutes		
Main unit weight	Approx.273g(batteries not included)		
Main unit size	130mmx95mmx52.5mm		
Main unit lifetime	10,000 times under normal use		
Battery life	Could be used for	Could be used for about 310 times when charge is full	
Accessories	Cuff, instruction manual, lithium battery, USB cable		
Operating environment	Temperature	5°C~40°C	
	Humidity	15%~93%RH	
	Air pressure	86kPa-106kPa	
Storage environment	Air pressure:86kPa-106kPa; Temperature:-20°C-55°C; Humidity:10%-93%RH; avoid crash,sun burn or rain during transportation		
Expected service life	Five years		
Software version	UA1.0		

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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.

Table 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		

FCC Statement

810

870

930

1720

1845

1970

2450

5240

5500

5785

800 - 96

1700 -

2570

5100 -

5800

Table 3

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Any Changes or modifications not expressly approved by the party responsible for compliance could void the user's authority to

Note: This equipment has been tested and found to comply with the limits for a Class B digital device, pursuant to part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:

-Reorient or relocate the receiving antenna.

-Increase the separation between the equipment and receiver. -Connect the equipment into an outlet on a circuit different from that to which the receiver is connected

-Consult the dealer or an experienced radio/TV technician for help.

FCC RF Exposure Information and Statement

When carrying the product or using it while worn on your body, either use an approved accessory such as a holster or otherwise maintain a distance of 5 mm from the body to ensure compliance with RF exposure uirements. Use of other accessories may not ensure compliance with FCC RF exposure guidelines.

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Arm Blood Pressure Monitor

Proper use of the unit

Memory-recall of measurements

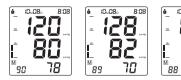
This blood pressure monitor automatically stores 2x90 sets measurements value, the oldest record will be replaced by the latest measurement value when there are more than 90 sets of memories in each user

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











Memory - clear of measurements

If you are sure that you want to permanently remove all stored memories. Press the button SET for 18 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.

Check IMFI details

After long pressing the MEM button for 5 seconds in the shutdown state, a bouncing bar "-" will appear in the upper right corner of the screen for about 2 seconds, and then the IMEI number of the device will be displayed on the screen. Press the START/STOP button to exit.



Exceptional Situation

Error indicators

■The following symbol will appear on the display when measuring abnormal

Cause	Correction	
Weak signal or	Wrap the cuff properly.	
suddenly	Remeasure with correct way.	
External strong	When near cell phone or other high radiant device , the measurement will be failed.	
disturbance	Keep quite and no chatting when measure.	
It appears error	Wrap the cuff properly.	
during the process of	Make sure that the air plug is properly inserted in the unit.	
inflating	Remeasure.	
Abnormal blood pressure	Repeat the measurement after relax for 30 mins , if get unusual readings for 3 times, please contact your doctor.	
Low battery	Connect the USB cable and charger to charge the battery.	
	Weak signal or pressure change suddenly External strong disturbance It appears error during the process of inflating Abnormal blood pressure	

Trouble removal

Problem	Check	Cause and solutions	
No power	Check the battery power	Charge the battery	
	Whether the plug insert	Insert into the air socket tightly	
No inflation	Whether the plug broken or leak	Change a new cuff	
Err and stop working	Whether move the arm when inflate	Keep the body peaceful	
Lif and stop working	Check if chatting when measured	Keep quite when measure	
Cuff look	Whether the cuff wrap too loose	Wrap the cuff tightly	
Cuff leak	Whether the cuff broken	Change a new cuff	
A Please contact the unit by yourself!	distributor if you can't solve the	problem, do not disassemble the	

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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 upper arm. ■ The unit satisfies the requirements of EN ISO 81060-1 Part 1 Noninvasive sphygmomanometers, EN 1060-3:1997+A2:2009 Non-invasive sphygmomanometers. 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 Five 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 ■ Monitor subjected to misuse, abuse, and neglect of these manual
- content, non-instructional purposes; unauthorized repair or modifications will be excluded from this warranty. ⚠ The device requires no calibration

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⚠ The device is not repairable and contains no user serviceable parts.

EMC Declaration

Immunity Test	EC 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 contact ±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
Surgel EC 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 70% 25/30 cycles Single phase: at 0 0% 250 cycle (50Hz)	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 0% 250 cycle (50Hz
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

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telli health

Manufacturer
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

Manufactured for:Telli Health LLC 66 West Flagler Street - Suite 900, Miami,FL,33130 USA



REV.00