材质：80克书写纸
成品尺寸：90*65mm
展开尺寸：360*260mm

Blood Pressure Monitor

Model Number: W06LT

USER MANUAL

Shenzhen Jamr Technology Co., Ltd.
A101-301, D101-201, Jamr Science & Technology Park, No. 2 Guiyuan Road, Guixiang Community, Guanlan Street, Longhua District, 518110 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

Published Date:2025/4/27
Version Number:1.0

CONTENTS

1 Introduction and Intended Use.....1
2 Important Information on Blood Pressure and its Measurement.....4
3 Components of your blood pressure monitor.....6
4 Using your Monitor for the First Time.....7
5 Measurement Procedure.....9
6 Care and Maintenance.....20
7 Warranty.....21
8 Certifications.....21
9 Technical Specifications.....22
10 FCC Statement.....23
11 EMC Declaration.....25

1 Introduction and Intended Use

The device is a fully automatic digital blood pressure measuring device using oscillometric technique to measure systolic and diastolic blood pressure as well as the pulse for adults that ages are more than 12 years old by wrapping around the wrist with cuff circumference ranging from 12.5 cm to 21.5 cm.The device can be used in medical facilities or at home, and only for indoor use.

The device is provide accurate blood pressure measurement values that are effective and suitable for clinical and home use.

Before using, please read this instruction manual carefully and then keep it in a safe place.

1.1 Remember...

• Only a health-care professional is qualified to interpret blood pressure measurements.
• This device is NOT intended to replace regular medical checkups.
• The device is intended for use by adults only and not intended for use on children and pregnant patient.The effectiveness has not been established in pregnant (including pre-eclampticl) patients.
•In cases of irregular heartbeat, measurements made with this instrument should only be evaluated after consultation with a physician.
• The products, including accessories, shall be processed in accordance with local regulations after reaching the life cycle.

• Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

1.2 Warnings and Precautions ⚠

Warning: The device is not suitable for use in the presence of flammable anesthetic mixtures with air or with oxygen or nitrous oxide.

Warning: If the patient is an intended operator, the functions of monitoring blood pressure and pulse rate can be safely used by patient. The routine clean can be performed by the patient.

Warning: Too frequent measurements can cause injury to the PATIENT due to blood flow interference.

Warning: Don't place the cuff over wound part.

Warning: Pressurization of the CUFF can temporarily cause loss of function of simultaneously used monitoring ME EQUIPMENT on the same limb.

Warning: Regularly checking the operation of the blood pressure monitor to ensure that it does not cause long-term damage to the patient's blood circulation.

Warning: Apply CUFF and its pressurization on the side of the patient's mastectomy or lymph node removal can cause injury.

Warning: Replacing a lithium battery or fuel cell with an inadequately trained person can lead to dangerous conditions (such as overtemperature, fire, or explosion).

Caution: The user must check that the equipment functions safely and see that it is in proper working condition before being used.

Caution: To avoid damaging the device, keep this unit away from children and pets.

Caution: The standard material used for the bladder is latex-free.

Caution: The device is intended to monitor, not to diagnose. Unusual values have to be always discussed with a physician. Under any circumstance, you should not alter the dosages of any drugs prescribed by a physician.

Caution: The device cannot be used to substitute the professional ECG monitor device for monitoring the frequency of heart beat.

Caution: This device can not be used together with HF surgical equipment.

Note: To obtain the greatest accuracy from your blood pressure monitor, it is recommended that the instrument be used within the specified temperature and the relative humidity, please see the Technical Specifications.

Note:The device can not be used in MRI environment.

Note: The cuff is defined as the applied part. The user should contact the manufacturer for assistance, if needed, replace, or maintaining the device.

Note: This device contains sensitive electronic components. Avoid strong electrical or electromagnetic fields in the direct vicinity of the device (e.g. mobile telephones,

microwave ovens) during use. These can lead to erratic results.

Note: Do not attempt to service or repair this device yourself. Should a malfunction occur, refer to local distributor or the manufacturer.

2 Important Information on Blood Pressure and its Measurement

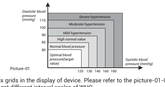
2.1 How does high or low blood pressure arise?

Your level of blood pressure is determined in the circulatory center of the brain and adjusts to a variety of situations through feedback from the nervous system. To adjust blood pressure, the strength and speed of the heart (Pulse), as well as the width of circulatory blood vessels is altered.Blood vessel width is controlled by fine muscles in the blood vessel walls.

Your level of arterial blood pressure changes periodically during heart activity: During the "blood ejection" (Systole) the value is highest (systolic blood pressure value). At the end of the heart's "rest period" (Diastole) pressure is lowest (diastolic blood pressure value).

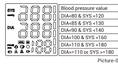
2.2 Which values are normal?

Please refer to the diagram below(Picture-01)



Picture-01

There are six grids in the display of device. Please refer to the picture-01 -01. Different grids represent different interval scales of WHO.

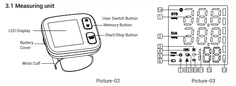


Blood pressure value	WHO grids in device	WHO Classification
DIA<80 & SYS <120	1	Optimal blood pressure
DIA<85 & SYS <130	2	Normal blood pressure
DIA<90 & SYS <140	3	High normal value
DIA<100 & SYS <160	4	Mild hypertension
DIA<110 & SYS <180	5	Moderate hypertension
DIA>=110 or SYS >=180	6	Severe hypertension

Picture-01-01

3 Components of your blood pressure monitor

3.1 Measuring unit



Picture-02

Picture-03

3.2 The symbols on the LED display

1. Systolic blood pressure
2. Diastolic blood pressure
3. Memory symbol
4. Average value symbol
5. Movement error symbol
6. Low battery symbol
7. User 1
8. User 2
9. Irregular heartbeat symbol
10. Mute symbol
11. Heartbeat symbol
12. Pulse display/ Memory number
13. WHO symbol
14. Bluetooth symbol

3.3 Features of Model W06LT

1. Average value function
2. Double users: 2 x 120 sets memory
3. WHO function
4. Irregular heartbeat checking
5. Auto power-off
6. Low battery display
7. Body movement detecting

4 Using your Monitor for the First Time

4.1 Battery Power checking

The battery is built-in chargeable Lithium battery.
Press the "START/STOP " button, if the symbol ( ) is blinking and the device speaks "battery low power, please recharge it". It means the battery power is low and need to be recharged.
During the charging process, the charging indicator on the display screen blinks. When the charging indicator stops blinking, it means the battery power is fully recharged.

4.2 System Settings

Before setting, ensure that the battery power is enough.

Setting the User ID(1 or 2)

With the unit off, press the ( ) button and then you can set the User ID by pressing the ( ) button again.

Setting the Volume

With the unit off, long press the ( ) button for more than 3s, and then you can start to set. When display with VOL is flashing, press the( )button to switch volume 1, volume 2, volume 3 or OFF. Press the ( ) button to confirm. After the setting is completed, the device switches off automatically and save the setting result.

5 Measurement Procedure

5.1 Before measurement

• Avoid eating and smoking as well as all forms of exertion directly before measurement. These factors influence the measurement result. Find time to relax by sitting in an armchair in a quiet atmosphere for about ten minutes before taking a measurement.
• Remove any garment that fits closely to your wrist.
• Always measure on the same wrist (normally left).

5.2 Fitting the Cuff

Please refer to picture-04

1. Remove all jewelry and watches. The palm of your hand should be facing you. Apply the cuff so that the display is facing you.
2. The distance between the cuff and the hand should be 1cm.
3. Secure the cuff with the hook and loop adhesive so that it lies comfortably and not too tight. It should be comfortable.
4. Lay your arm on a table with your palm upwards. Support your arm with a cushion so that the cuff rests at about the same height as the heart. Remain still for 2 minutes, sitting quietly, before beginning a measurement.

Picture-04

Operate via the App on smart phone with Bluetooth

1. Install the App from Google play store or Apple app store. Open Bluetooth on smart phone, and then turn on the App.
2. Bluetooth pairing
Turn on the device and the Bluetooth symbol ( ) will flash, Then operate bluetooth pairing according to the Settings on the APP. The Bluetooth symbol ( ) will stop flashing after the Bluetooth pairing is successful.
3. Bluetooth will be automatically searched and connected when it is powered on. After the Bluetooth connection is successful, the Bluetooth symbol ( ) will stop flashing and the measurement data will be uploaded to the APP.
Note:Devices that have been successfully paired will save the pairing information and do not need to be paired again.
It is recommended to connect the APP through Bluetooth before each measurement and then start the measurement.

5.3 Measure Procedure

The device is designed to take measurements and store the measurement values in memory for two people using User ID 1 and User ID 2.
Refer to picture-05
1. Sit comfortably in a chair with your feet flat on the floor.
2. Select your User ID (1 or 2).
Stretch your arm forward on the desk and keep relaxing, make sure the palm of hand is upturned. Make sure wrist is in correct position, to avoid body movement. Sit still and do not talk or move during the measurement.
After the cuff has been appropriately positioned on the wrist and connected to the blood pressure monitor, the measurement can begin:

Operate on the device

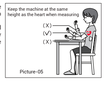
1. Press the START/STOP button. The pump begins to inflate the cuff. In the display, the increasing cuff pressure is continually displayed.
2. After automatically reaching an individual pressure, the pump stops and the pressure slowly falls. The cuff pressure is displayed during the measurement.
3. When the device has detected your pulse, the heart symbol in the display begins to blink.
4. When the measurement has been concluded, the measured systolic and diastolic blood pressure values, as well as the pulse will be displayed.
5. The measurement results are displayed until you switch the device off. If no button is pressed for 60 seconds, the device switches off automatically.

6. Movement error symbol ( )
The Movement Error Symbol ( ) is displayed if you move your body during the measurement. Please remove the cuff, and wait 2-3 minutes. Reapply the cuff and take another measurement.

NOTE:
Patient Position:
1. Comfortably seated
2. Legs uncrossed
3. Feet flat on the floor
4. Back and arm supported
5. Middle of the CUFF at the level of the right atrium of the heart

Recommended Use Methods
1. Recommendation that the PATIENT relax as much as possible and not talk during the measurement PROCEDURE
2. Recommendation that 5 min should elapse before the first reading is taken
3. Any reading can be affected by the measurement site, the position of the PATIENT, exercise, or the PATIENT'S physiologic condition
4. Performance of the AUTOMATED SPHYGMOMANOMETER can be affected by extremes of temperature, humidity and altitude.

5. To stop the inflation or measurement, push the START/STOP button. The monitor will stop inflating, start deflating, and will turn off.
6. After the monitor has detected your blood pressure and pulse rate, the cuff automatically deflates. Your blood pressure and pulse rate are displayed.



Picture-05

5.4 Irregular Heartbeat Detector

This symbol ( ) indicates that certain pulse irregularities were detected during the measurement.
In this case, the result may deviate from your normal basal blood pressure -- repeat the measurement.
Information for the doctor on frequent appearance of the Irregular Heartbeat Symbol.
This instrument is an oscillometric blood pressure monitor device that also analyzes pulse frequency during measurement. The instrument is clinically tested.
If pulse irregularities occur during measurement, the irregular heartbeat symbol is displayed after the measurement. If the symbol appears more frequently (e.g. several times per week on measurements performed daily) or if it suddenly appears more often than usual, we recommend the patient to seek medical advice. The instrument does not replace a cardiac examination, but serves to detect pulse irregularities at an early stage.

5.5 Error Indicates

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

SYMBOL	CAUSE	CORRECTION
No display appears	Weak battery	Please charge in time.
E1	Sensor abnormal	Check if the pump is working or not. If it is working, then the problem is sensor abnormal. Please send it to the local distributor.
E2	Monitor could not detect pulse wave or insufficient pulse wave data	Tie the wrist cuff correctly and make the measurement again. If the error is still displayed, please send it to local distributor.
E3	Measurement results is abnormal or out the measurable range of blood pressure	Please keep quiet and measure again
E4	Too loose cuff or air leakage	Correct it and make the measurement again
E5	The air tube is crimped or the wrist cuff is tied too tight	Correct it and make the measurement again
E6	The sensor is sensing great fluctuation in the pressure	Please keep quiet and don't move
E7	The pressure that the sensor sensing is over the limit	Start the measurement again if the error is still displayed, please send it to local distributor
E8	The demarcation is incorrect or the device has not been demarcated	Please send back to the local distributor

HI	The pulse rate exceeds the upper limit (>199 per minute)	Beyond the measurement range, normal reminder	
LO	The pulse rate is less than the lower limit (<40 per minute)	Beyond the measurement range, normal reminder	

Trouble removal

Problem	Check	Cause and solutions
No power	Check the battery power	Charge the battery
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 shaking 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

⚠ Please contact the distributor if you can't solve the problem, do not disassemble the unit by yourself!

SYMBOL DESCRIPTIONS

The following symbols may appear in this manual, on the Digital Blood Pressure Monitor W06LT, or on it's accessories. Some of the symbols represent standards and compliances associated with the Digital Blood Pressure Monitor W06LT and its use.

	Date of manufacture.
	Manufacturer.
SN	Specifies serial number.
	Type BF applied part.
	Direct current.
	DISPOSAL: Do not dispose this product as unsorted municipal waste. Collection of such waste separately for special treatment is necessary.
	Follow instructions for use.
IP22	The degree of avoid ingress of water or particulate matter into ME equipment
	MR unsafe
MD	Medical device

	Put up		Handle gently
	Keep dry		Temperature range
	Fragile		Avoid the sun
No Sterilize requirement			
Not category AP / AG equipment			
Mode of operation: continuous			

5.6 Memory

Each unit stores 120 sets measurements for 2 users, totally 240 sets (User 1 and 2). Measurements for each user are stored separately. Be certain that you are viewing the measurements for the correct user.

View the memory

With the unit off, press the ( ) button. The monitor will display User ID and an average value of the last 3 times measurements stored in the unit. (If measurements are less than 3 sets, directly display the first set).

Each time you press the ( ) button, it will display the memory value from the latest to the oldest in turn.
Each time you press the( ) button, it will display the memory value from the oldest to the latest in turn.

Delete memory

In average value memory viewing mode,the average value symbol ( ) is being displayed, long press the ( ) button for 3 seconds, then it will delete all measurements for the current user.
In single set memory viewing mode, long press the ( ) button for 3 seconds, then it will delete only a set measurement being displayed.

Note:

If you decide to delete the all memory, please keep the memory in another way, incase you need it some days later.

5.7 Discontinuing a Measurement

If it is necessary to interrupt a blood pressure measurement for any reason (e.g the patient feels unwell), the Start/Stop button can be pressed at any time. The device then immediately lowers the cuff pressure automatically.

6 Care and Maintenance

Wash hands after each time measurement.
If one device is used by different patients, wash hands before and after each use.
1. Do not expose the device to either extreme temperatures, humidity, dust or direct sunlight.
2. The cuff contains a sensitive air-tight bladder. Handle this cuff carefully and avoid all types of stress through twisting or buckling.
3. Clean the device with a soft, dry cloth. Do not use gas, thinners or similar solvents. Spots on the cuff can be removed carefully with a damp cloth and soapsuds. The cuff with bladder must not be washed in a dishwasher, clothes washer, or submerged in water.
4. Keep the device away from sharp edges.
5. Do not drop the monitor or treat it roughly in any way. Avoid strong vibrations.
6. Never open the monitor! This invalidates the manufacturer's warranty.
7. Batteries and electronic instruments must be disposed of in accordance with the locally applicable regulations, not with domestic waste.

6.1 Accuracy test

Sensitive measuring devices must be checked for accuracy from time to time. We recommend a periodical inspection of your unit by an authorized dealer every 1 year. Please turn to local distributor or the manufacturer.

7 Warranty

Your blood pressure monitor is guaranteed for 2 years against manufacturers' defects for the original purchaser only, from date of purchase. The warranty does not apply to damage caused by improper handling, accidents, unprofessional use, not following the operating instructions or alterations made to the instrument by third parties.
Warranty only applies to the main device and its cuff. All other accessories are not covered by warranty.
There are no user serviceable parts inside. Battery is not covered by the warranty.
Note: According to international standards, your monitor should be checked for accuracy every year.

8 Certifications

Device standard:
This device is manufactured to meet the blood pressure monitors:
IEC 80601-2-30 / IEC60601-1-11 / IEC60601-1
Electromagnetic compatibility:
Device fulfills the stipulations of the International standard IEC60601-1-2
The device was clinically investigated and the safety and efficacy is meet the requirement of ISO 81060-2. If you need to acquire a copy of the summary of the Clinical Investigation, please contact the manufacturer.

9. Technical Specifications

Model: W06LT
Weight: 102g (Battery is included)
Display: 45×45mm (1.77"×1.77") LED Digital Display
Size: 76(W)×66.0(+23.0) mm (3.11"(W)×2.6"(L)×0.91"(H))
Accessories: 1×Main Device, 1×Cuff, 1×Users manual,
Operating Conditions: Temperature: 5 ~ to 40 ; Humidity: 15% to 90% RH.
Storage And Shipping Conditions: Temperature: -20 to 60 ; Humidity: 10%RH~93%RH
Atmospheric pressure range: 70kPa~106kPa
Measuring method: Oscillometric
Pressure sensor: Resistive
Measuring range: DIA: 40-130mmHg; SYS: 60-230mmHg
Pulse: 40-199 beats/min
Cuff pressure display range:0-299mmHg
Memory: Automatically stores the last 120 measurements for 2 users (total 240)
Measuring resolution: 1 mmHg
Accuracy: Pressure within ± 3 mmHg / pulse ± 5 % of the reading
Power source: Built-in high capacity lithium battery - 300mAh
Cuff: 4.92"-8.46" (12.5-21.5cm)
Users: Adult
Expected service life of the device: 5 years
Technical alterations reserved!

10. FCC Statement

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.
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.

Caution: Any changes or modifications not expressly approved by the party responsible for compliance could void the user's authority to operate the equipment.
This equipment complies with RF radiation exposure limits set forth for an uncontrolled environment.
This transmitter must not be co - located or operating in conjunction with any other antenna or transmitter.
The device has been evaluated to meet general RF exposure requirement. The device can be used in portable exposure condition without restriction.

11 EMC Declaration

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 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 (W06LT), 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

Table 2

Guidance and manufacturer's declaration - electromagnetic 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 contact ±2 kV, ±4 kV, ±8 kV, ±15 kV air
Electrical fast transient/burst IEC 61000-4-4	Power supply lines: ±2 kV Input/output lines: ±1 kV 100 kHz repetition frequency	Not application
Surge IEC 61000-4-5	Line(s) to line(s): ±1 kV line(s) to earth: ±2 kV	Not application
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	Not application
Power frequency magnetic field IEC 61000-4-8	3A/m 50Hz/60Hz	30 A/m 50Hz/60Hz
Conducted RF IEC61000-4-6	150kHz to 80MHz 3Vrms 6Vrms (in ISM and amateur radio bands) 80% AM at 1kHz	Not application
Radiated RF IEC 61000-4-3	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
Proximity magnetic fields IEC 61000-4-39	30 kHz: 8A/m 134.2 kHz: 65A/m 13.56 MHz: 7.5A/m	30 kHz: 8A/m 134.2 kHz: 65A/m 13.56 MHz: 7.5A/m

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

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).

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	Not application
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Not application

Table 3

Guidance and manufacturer's declaration - electromagnetic Immunity

	Test Frequency (MHz)	Band[a] (MHz)	Service [a]	Modulation[b]	IMMUNITY TEST LEVEL (V/m)
Radiated RF (Test specifications for enclosure port IMMUNITY RF wireless communication equipment)	385	380- 390	TETRA 400	Pulse modulation[b] 18 Hz	27
	450	430- 470	GMRS 460, FRS 460	FM [c] ±5 kHz deviation 1 kHz sine	28
	710	704- 787	LTE Band 13, 17	Pulse Modulation[b] 217 Hz	9
	745				
	780				
	810	800- 960	GSM 800/900, TETRA 800, iDEN 820, CDMA 850, LTE Band 5		28
	870				
	930				
	1720	1700- 1990	GSM 1800; CDMA 1900; GSM 1900; DECT; LTE Band 1, 3, 4, 25; UMTS	Pulse Modulation[b] 217 Hz	28
	1845				
	1970				
	2450	2400- 2570	Bluetooth,WLAN, 802.11 b/g/n, RFID 2450, LTE Band 7	Pulse Modulation[b] 217 Hz	28

5240	WLAN 802.11 a/n	Pulse Modulation[b] 217 Hz	9
5500			
5785			

5100-5800

a) If necessary to achieve the IMMUNITY TEST LEVEL, the distance between the transmitting antenna and the ME EQUIPMENT or ME SYSTEM may be reduced to 1 m. The 1 m test distance is permitted by IEC 61000-4-3.

a) For some services, only the uplink frequencies are included.
b) The carrier shall be modulated using a 50 % duty cycle square wave signal.
NOTE In cases where pulse modulation is used, the modulation shall exceed a 50 % duty cycle square wave signal at 18 Hz. While it does not represent actual modulation, it would be worst case.

Table 4

Guidance and manufacturer's declaration - electromagnetic Immunity

Test frequency	Modulation	IMMUNITY TEST LEVEL (A/m)
30 kHz[a]	CW	8
134.2 kHz	Pulse modulation a) [b] 2.1 kHz	65 [c]
13.56 MHz	Pulse modulation a) 50 kHz	7.5[c]

a) This test is applicable only to ME EQUIPMENT and ME SYSTEMS intended for use in the HOME HEALTHCARE ENVIRONMENT.
a) The carrier shall be modulated using a 50% duty cycle square wave signal.
b) c) m.s., before modulation is applied.