

Instruction for use



B.Well

Caring for everyone

PRO-36

Automatic blood pressure monitor

High-accuracy
measurements

ENG



 PRO

 MED

 TECHNO

Model PRO-36

Automatic Upper Arm Blood Pressure Monitor

Instruction for Use

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1. INTRODUCTION

Thank you for purchasing the B.Well “talking” upper arm blood pressure monitor PRO-36. Designed for convenient and easy operation, this device provides fast and reliable measurement of systolic and diastolic blood pressure as well as heart rate using the oscillometric measurement method.

Your blood pressure is an important parameter that can be used to monitor your health. High blood pressure (hypertension) is a serious health problem that is frequently found in the modern world. This device enables you to monitor your blood pressure regularly.

The PRO-36 is a voice-guided fully automatic, digital, upper arm blood pressure measuring device.

- Up-to-date **IntellectClassic** technology uses oscillometric measurement during deflation for quick, precise and painless result.
- The Pulse Arrhythmia Detection technology with an audible sound signal which warns about the malfunctions of normal rate and periodicity of systole during the measurement
- Big 3-line LCD screen
- One convenient button with backlight
- Last measurement memory
- Fan-shape anatomic cuff for arm, washable
- Battery life indicator
- Automatic switch off
- The possibility to use mains adapter
- This device is easy to use and has been proven in clinical studies to provide excellent accuracy.
- Before using the PRO-36, read this instruction manual carefully and keep it in a safe place. For further questions on the subject of blood-pressure and its measurement, please contact your doctor.
- Voice-guided with adjustable volume knob

2. IMPORTANT INFORMATION ON THE SUBJECT OF BLOOD-PRESSURE AND ITS MEASUREMENT

2.1. Normal blood pressure fluctuation

All physical activity, excitement, stress, eating, drinking, smoking, body posture and many other activities or factors (including taking a blood pressure measurement) will influence blood pressure value. Because of this, it is mostly unusual to obtain identical multiple blood pressure readings.

Blood pressure fluctuates continually day and night. The highest value usually appears in the daytime and lowest one usually at midnight. Typically, the value begins to increase at around 3:00 AM, and reaches to highest level in the daytime while most people are awake and active.

Considering the above information, it is recommended that you measure your blood pressure at approximately the same time each day. Too frequent measurements may cause injury due to blood flow interference, please always relax a minimum moment of 1 to 1.5 minutes between measurements to allow the blood circulation in your arm to recover. It is rare that you obtain identical blood pressure readings each time.

The device contains sensitive electronic components (Microcomputer). Therefore, avoid strong electrical or electromagnetic fields in the direct vicinity of the device (e.g. mobile telephones, microwave cookers). These can lead to temporary impairment of the measuring accuracy.

2.2. Classification of blood pressure values

Blood pressure is too high if at rest, the diastolic pressure is above 90mmHg and/or the systolic blood-pressure is over 160mmHg. In this case, please consult your doctor immediately. Long-term values at this level endanger your health due to the associated advancing damage to the blood vessels in your body.

Should the systolic blood-pressure values lie between 140mmHg and 160mmHg and/or the diastolic blood-pressure values lie between 90mmHg and 100mmHg, likewise, please consult your doctor. Furthermore, regular self-checks will be necessary.

With blood-pressure values that are too low, i.e. systolic values under 100mmHg and/or diastolic values under 60mmHg, likewise, please consult your doctor.

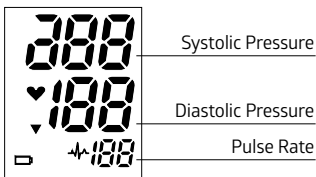
Even with normal blood-pressure values, a regular self-check with your blood-pressure monitor is recommended. In this way you can detect possible changes in your values early and react appropriately.

If you are undergoing medical treatment to control your blood pressure, please keep a record of the level of your blood pressure by carrying out regular self-measurements at specific times of the day. Show these values to your doctor. Never use the results of your measurements to alter independently the drug doses prescribed by your doctor.

Table for classifying blood-pressure values (unit: mmHg) according to European Society of Hypertension (ESH)

Range	Systolic blood pressure	Diastolic blood pressure	Measures
Grade 3: severe hypertension	Higher or equal to 180	Higher or equal to 110	Urgently seek medical advice!
Grade 2: moderate hypertension	160-179	100-109	Consult your doctor immediately
Grade 1: mild hypertension	140-159	90-99	Consult your doctor
High normal	130-139	85-89	Consult your doctor
Normal	Lower than 130	Lower than 85	Self-check
Optimal	Lower than 120	Lower than 80	Self-check

3. CONTENTS AND DISPLAY INDICATORS



- ♥ Heart beat symbol
- ▼ Zero-seeking symbol
- 🔋 Low battery symbol
- 📶 Pulse Arrhythmia Detection symbol





4. INTENDED USE

The digital automatic blood pressure monitor is for use by medical professionals or at home and is a non-invasive blood pressure measurement system intended to measure the diastolic and systolic blood pressures and pulse rate of an adult individual by using a non-invasive technique in which an inflatable cuff is wrapped around the upper arm. The cuff circumference is limited to 22cm-48cm.

5. CONTRAINDICATION

It is inappropriate for people with serious arrhythmia to use the digital automatic blood pressure monitor.




6. PRECAUTIONS


1. Read all of the information in the operation guide and any other literature in the box before operating the unit.
2. Stay still, calm and rest for 5 minutes before blood pressure measurement.
3. The cuff should be placed at the same level as your heart.
4. During measurement, neither speak nor move your body and arm.
5. Measuring on left arm for each measurement.
6. Please always relax a minimum moment of 1 to 1.5 minutes between measurements to allow the blood circulation in your arm to recover. Prolonged over-inflation (cuff pressure exceed 300 mmHg or maintained above 15 mmHg for longer than 3 minutes) of the bladder may cause ecchymoma of your arm.
7. Consult your physician if you have any doubt about below cases:
 - 1) The application of the cuff over a wound or inflammation diseases;
 - 2) The application of the cuff on any limb where intravascular access or therapy, or an arterio-venous (A-V) shunt, is present;
 - 3) The application of the cuff on the arm on the side of a mastectomy;
 - 4) Simultaneously used with other monitoring medical equipments on the same limb;
 - 5) Need to check the blood circulation of the user.
8.  This digital automatic blood pressure monitor is designed for adults and should never be used on infants or young children. Consult your physician or other health care professionals before use on older children.
9. Do not use this unit in a moving vehicle, This may result in erroneous measurement.
10. Blood pressure measurements determined by this monitor are equivalent to those obtained by a trained observer using the cuff/stethoscope auscultation method, within the limits prescribed by the American National Standard Institute, Electronic or automated sphygmomanometers.
11. Information regarding potential electromagnetic or other interference between the blood pressure monitor and other devices together with advice regarding avoidance of such interference please see part ELECTROMAGNETIC COMPATIBILITY INFORMATION.
12. If certain pulse irregularity caused by common arrhythmias is detected in the procedure of blood pressure measurement, a signal of  will be displayed. Under this condition, the digital automatic blood pressure monitor can keep function, but the results may not be accurate, it's suggested that you consult with your physician for accurate assessment.
There are 2 conditions under which the signal of Pulse Arrhythmia Detection will be displayed:
 - 1) The coefficient of variation (CV) of pulse period >25%.
 - 2) The difference of adjacent pulse period $\geq 0.14s$, and the number of such pulse takes more than 53 percentage of the total number of pulse.
13. Please do not use the cuff other than supplied by the manufacturer, otherwise it may bring biocompatible hazard and might result in measurement error.

14. ⚠ The monitor might not meet its performance specifications or cause safety hazard if stored or used outside the specified temperature and humidity ranges in specifications.
15. ⚠ Please do not share the cuff with other infective person to avoid cross-infection.
16. Please note that changes or modifications not expressly approved by the party responsible for compliance could void the user's authority to operate the equipment.
17. 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.
18. Medical AC adapter which output is DC 6.0V 600mA and complied with IEC 60601-1/EN 60601-1/UL 60601-1 and IEC 60601-1-2/EN 60601-1-2/UL 60601-1-2 is suitable for this monitor. Please note that the monitor jack size: hole \varnothing 5.5mm, center pin \varnothing 2.0mm. Please pay attention to polarity.

7. SETUP AND OPERATING PROCEDURES

7.1. Battery loading


- a. Open battery cover at the back of the monitor.
- b. Load four "AAA" size batteries. Please pay attention to polarity.
- c. Close the battery cover.
- d. Once you install the batteries or turn off the monitor, the LCD does not display anything. Now the monitor is in Off Mode.
 - ⚠ If the LCD display battery symbol  while the monitor is on, the batteries are running out.
 - ⚠ If the batteries are run out, battery symbol  will blink for 10 seconds. Then the monitor will always display battery symbol  and cannot open. Please replace all batteries with new ones.
 - ⚠ Rechargeable batteries are not suitable for this monitor.
 - ⚠ Remove the batteries if the monitor will not be used for a month or more to avoid relevant damage of battery leakage.
 - ⚠ Avoid the battery fluid to get in your eyes. If it should get in your eyes, immediately rinse with plenty of clean water and contact a physician.

 *The monitor, the batteries and the cuff, must be disposed of according to local regulations at the end of their usage.*

7.2. Using a mains adapter

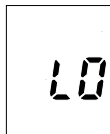
- 1) Plug the mains adaptor into a 100-240 V, 50/60Hz power socket.
- 2) Plug the DIN plug into the socket at the right side of the instrument.

No power is taken from the batteries while the mains adaptor is connected to the instrument.

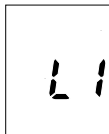
 **Warning** If You need mains adapter, You may purchase it separately. Use only the mains adaptor AD-155. The use of any other adapters can make your warranty void.

7.3. Voice setting

- a. Voice language setting: In turn-off mode, you can select the voice language by keeping on press the "START" button. Now LCD blink "L0", "L1", "L2" ... "Ln" (n means the number of language) circularly, see Picture 2 & Picture 2-1. "L0" represents closing voice function, "L1" represents language 1, "L2" represents language 2, ..., "Ln" represents language n, and so on. You can select the wanted language by releasing "START" button when display the corresponding language code.
- b. Voice volume setting: Once you have selected a language, while the monitor is speaking something, you can roll the Voice Knob to setting the voice volume.




Picture 2



Picture 2-1

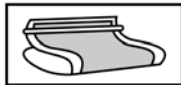
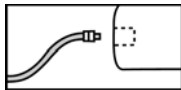
7.4. Connecting the cuff to the monitor

Insert the Air Tube Plug firmly into the Air Tube Socket on the side of the Monitor. Make certain that the Plug is completely inserted in order to prevent air leakage during use.

 Avoid compression or restriction of the connection tubing during measurement, which may cause inflation error, or harmful injury due to continuous cuff pressure.

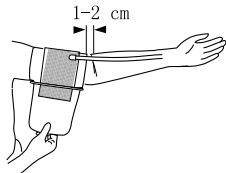
7.5. Applying the cuff

- a. Pulling the cuff end through the medal loop (the cuff is packaged like this already), turn it outward (away from your body) and tighten it and close the Velcro fastener.
- b. Place the cuff around a bare left arm 1-2cm above the elbow joint.



Instruction for use

- While seated, place palm upside in front of you on a flat surface such as a desk or table. Position the air tube in the middle of your arm in line with your middle finger.
- The cuff should fit comfortably, yet snugly around your arm. You should be able to insert one finger between your arm and the cuff.

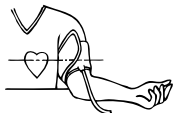


- Note:**
- Please refer to the cuff circumference range in "SPECIFICATIONS" to make sure that the appropriate cuff is used.
 - Measure on the left arm each time.
 - Do not move your arm, body, or the monitor and do not move the rubber tube during measurement.
 - Stay still, calm for 5 minutes before blood pressure measurement.
 - Please keep the cuff clean. If the cuff becomes dirty, remove it from the monitor and clear it by hand in a mild detergent, then rinse it thoroughly in cold water. Never dry the cuff in clothes dryer or iron it. Clean the cuff after the usage of every 200 times is recommended.
 - Do not place the cuff around your arm if the arm has any inflammation, acute diseases, infections skin wounds.

7.6. Carrying out a measurement

Before the measurement:

- Avoid eating, smoking as well as all forms of exertion directly before the measurement. All these factors influence the measurement result. Try and find time to relax by sitting in an armchair in a quiet atmosphere for about ten minutes before the measurement.
- Remove any garment that fits closely to your upper arm.
- Measure always on the same arm (normally left).
- Attempt to carry out the measurements regularly at the same time of day, since the blood-pressure changes during the course of the day.

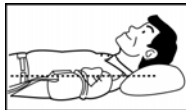


Sitting Comfortably Measurement

- Be seated with your feet flat on the floor, and don't cross your legs.
- Place palm upside in front of you on a flat surface such as a desk or table.
- The middle of the cuff should be at the level of the right atrium of the heart.

Lying Down Measurement

- Lie on your back.
- Place your left arm straight along your side with your palm upside.
- The cuff should be placed at the same level as your heart.

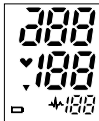


Common sources of error:

- ❗ **Note:** Comparable blood-pressure measurements always require the same conditions! These are normally always quiet conditions.
- All efforts by the patient to support the arm can increase the blood-pressure. Make sure you are in a comfortable, relaxed position and do not activate any of the muscles in the measurement arm during the measurement. Use a cushion for support if necessary.
 - If the arm artery lies considerably lower (higher) than the heart, an erroneously higher (lower) blood-pressure will be measured! (Each 15cm difference in height results in a measurement error of 10mmHg)
 - Cuffs that are too narrow or too short result in false measurement values. Selecting the correct cuff is of extraordinary importance. The cuff size is dependent upon the circumference of the arm (measured in the centre). The permissible range is printed on the cuff.
- ❗ **Note:** only use clinically approved Original-Cuffs!
- A loose cuff or a sideways protruding air-pocket causes false measurement values.
 - With repeated measurements, blood accumulates in the respective arm, which can lead to false results. Correctly executed blood-pressure measurements should therefore first be repeated after a 1 minute pause.

7.7. Taking your blood pressure reading

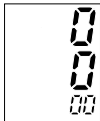
- After applying the cuff and your body is in a comfortable position, press the "START" button. The device reminds you: "Attention – the measurement is starting – please relax, do not move or talk" A beep is heard and all display characters are shown for self-test. See picture 3. Please contact the service center if segment is missing.
- The LCD will momentarily display the last measurement stored in the memory as well the device announces by 4 times the last memory reading. See picture 3-1. If the monitor has no measurement stored in the memory, the LCD will display show "0" for blood pressure and pulse rate. See picture 3-2.



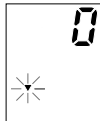
Picture 3



Picture 3-1



Picture 3-2



Picture 3-3



Picture 3-4

- Then the monitor starts to seek zero pressure. See picture 3-3.
- The monitor inflates the cuff until sufficient pressure has built up for a measurement. Then the monitor slowly releases air from the cuff and carries out the measurement. Finally the blood pressure and pulse rate will be calculated and displayed on the LCD screen and device automatically voices out your measured values and status, by 4 times.. Irregular heartbeat symbol (if any) will blink. "Additionally, pulse arrhythmia was detected..." will sound. See picture 3-4.



- e. After measurement, the monitor will turn off automatically after 1 minute of no operation. Alternatively, you can press the "START" button to turn off the monitor manually.
- f. During measurement, you can press the "START" button to turn off the monitor manually. The device then immediately lowers the cuff-pressure automatically, and you will hear at the time "B.Well wishes you all the best – stay healthy!"

① **Note:** Please consult a health care professional for interpretation of pressure measurements.


① **Note:** The monitor can memorize the last result. If you change the batteries, the last result may be lost.

7.8. Pulse Arrhythmia Detection

Appearance of the Arrhythmia indicator

The appearance of the symbol  signifies that a certain pulse irregularity was detected during the measurement. The result can vary from your normal blood pressure. As a rule this is not a cause for concern; however, 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 you inform your doctor. Please show your doctor the following explanation:

Information for the doctor on frequent appearance of the Arrhythmia indicator

The device is an oscillometric blood pressure measuring device that also analyses the pulse frequency during measurement. The accuracy of this device has been clinically validated. If pulse irregularities occur during measurement, the arrhythmia 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 device does not replace a cardiac examination, but serves to detect pulse irregularities at an early stage.

7.9. Technical alarm description

The monitor will show 'HI' or 'Lo' as technical alarm on LCD with no delay if the determined blood pressure (systolic or diastolic) is outside the rated range specified in part SPECIFICACIONES. In this case, you should consult a physician or check if your operation violated the instructions.


The technical alarm condition (outside the rated range) is preset in the factory and cannot be adjusted or inactivated. This alarm condition is assigned as low priority according to IEC 60601-1-8.

The technical alarm is non-latching and need no reset. The signal displayed on LCD will disappear automatically after about 8 seconds in the meantime, device reads out the possible causes..

7.10. Troubleshooting (1)

PROBLEM	POSSIBLE CAUSE	SOLUTION
LCD Display shows abnormal result	The cuff position was not correct or it was not properly tightened	Apply the cuff correctly and try again
	Body posture was not correct during testing	Review the "BODY POSTURE DURING MEASUREMENT" sections of the instructions and re-test.
	Speaking, arm or body movement, angry, excited or nervous during testing	Re-test when calm and without speaking or moving during the test
	Irregular heartbeat (arrhythmia)	It is inappropriate for people with serious arrhythmia to use this blood pressure monitor.

7.11. Troubleshooting (2)

PROBLEM	POSSIBLE CAUSE	SOLUTION
LCD shows low battery symbol 	Low Battery	Change the batteries
LCD shows "Er 0"	Pressure system is unstable before measurement	Don't move and try again.
LCD shows "Er 1"	Fail to detect systolic pressure	
LCD shows "Er 2"	Fail to detect diastolic pressure	
LCD shows "Er 3"	Pneumatic system blocked or cuff is too tight during inflation	Apply the cuff correctly and try again
LCD shows "Er 4"	Pneumatic system leakage or cuff is too loose during inflation	

LCD shows "Er 5"	Cuff pressure above 300mmHg	Measure again after five minutes. If the monitor is still abnormal, please contact the local distributor or the factory.
LCD shows "Er 6"	More than 3 minutes with cuff pressure above 15 mmHg	
LCD shows "Er 7"	EEPROM accessing error	
LCD shows "Er 8"	Device parameter checking error	
LCD shows "Er A"	Pressure sensor parameter error	
No response when you press button or load battery.	Incorrect operation or strong electromagnetic interference.	Take out batteries for five minutes, and then reinstall all batteries.

8. MAINTENANCE

1. ⚠ Do not drop this monitor or subject it to strong impact.
2. ⚠ Avoid high temperature and solarization. Do not immerse the monitor in water as this will result in damage to the monitor.
3. If this monitor is stored near freezing, allow it to acclimate to room temperature before use.
4. ⚠ Do not attempt to disassemble this monitor.
5. If you do not use the monitor for a long time, please remove the batteries.
6. It is recommended the performance should be checked every 2 years or after repair. Please contact the service center.
7. Clean the monitor with a dry, soft cloth or a soft cloth squeezed well after moistened with water, diluted disinfectant alcohol, or diluted detergent.
8. No component can be maintained by user in the monitor. The circuit diagrams, component part lists, descriptions, calibration instructions, or other information which will assist the user's appropriately qualified technical personnel to repair those parts of equipment which are designated repairably can be supplied.
9. The monitor can maintain the safety and performance characteristics for a minimum of 10,000 measurements or three years, and the cuff integrity is maintained after 1,000 open-close cycles of the closure.
10. It is recommended the cuff should be disinfected 2 times every week if needed (For example, in hospital or in clinique). Wipe the inner side (the side contacts skin) of the cuff by a soft cloth squeezed after moistened with Ethyl alcohol (75-90%), then dry the cuff by airing.

9. SPECIFICATIONS

1. Product name: Blood Pressure Monitor
2. Model: PRO-36
3. Classification: Internally powered, Type BF applied part, IPX0, No AP or APG, Continuous operation
4. Machine size: 87mm x 122mm x 53mm (3 7/16" x 4 13/16" x 2 3/32")
5. Cuff circumference: 22cm~42cm (8 21/32"~16 17/32") or 22cm~32cm (8 21/32"~ 12 19/32"), 30cm~42cm (11 13/16"~ 16 17/32") (Optional), 42cm~48cm (16 17/32"~18 29/32") (Optional)
6. Weight: approx. 195g (6 7/8 oz.) (exclude batteries and cuff)
7. Measuring method: oscillometric method, automatic air inflation and measurement
8. Memory volume: only the last measurement stored in the memory
9. Power source: DC6V \equiv 600mA, batteries: 4 \times 1.5V \equiv SIZE AAA
Mains adapter (optional)
10. Measurement range:
Cuff pressure: 0-300mmHg
Systolic: 60-260mmHg
Diastolic: 40-199mmHg
Pulse rate: 40-180 beats/minute
11. Accuracy:
Pressure: \pm 3mmHg
Pulse rate: \pm 5%
12. Environmental temperature for operation: 10°C~40°C(50°F~104°F)
13. Environmental humidity for operation: \leq 85% RH
14. Environmental temperature for storage and transport: -20°C~50°C(-4°F~122°F)
15. Environmental humidity for storage and transport: \leq 85% RH
16. Environmental pressure: 80KPa-105KPa
17. Battery life: Approx.270 times
18. List of all components belonging to the pressure measuring system, including accessories: Pump, Valve, LCD, Cuff, Sensor.

Note: These specifications are subject to change without notice.

10. APPLIED STANDARDS

The digital automatic blood pressure monitor corresponds to the below standards:

IEC 60601-1:2005/EN 60601-1:2006/AC:2010 (Medical electrical equipment – Part 1: General requirements for basic safety and essential performance),

IEC60601-1-2:2007/EN 60601-1-2:2007 /AC:2010 (Medical electrical equipment – Part 1-2: General requirements for basic safety and essential performance – Collateral standard: Electromagnetic compatibility – Requirements and tests),

IEC 80601-2-30 : 2009+Cor.2010 (Medical electrical equipment – Part 2-30: Particular requirements for the basic safety and essential performance of automated non-invasive sphygmomanometers),

EN 1060-1: 1995 + A1: 2002 + A2: 2009 (Non-invasive sphygmomanometers – Part 1: General requirements),

EN 1060-3: 1997 + A1: 2005 + A2: 2009 (Non-invasive sphygmomanometers – Part 3: Supplementary requirements for electro-mechanical blood pressure measuring systems).

11. SYMBOL INFORMATION



THE OPERATION GUIDE MUST BE READ
(The sign background colour: blue.The sign graphical symbol: white)



WARNING



TYPE BF APPLIED PARTS (The cuff is type BF applied part)



ENVIRONMENT PROTECTION – Waste electrical products should not be disposed of with household waste. Please recycle where facilities exist. Check with your local Authority or retailer for recycling advice.



MANUFACTURER'S NAME



ARTICLE NUMBER



SERIAL NUMBER



0044

CE mark (0044) COMPILES WITH MDD93/42/EEC REQUIREMENTS



POLARITY OF D.C. POWER CONNECTOR



OPERATING CONDITION, TEMPERATURE 10°C ~ 40°C



STORAGE CONDITION, TEMPERATURE -20°C ~ 50°C



KEEP DRY

12. WARRANTY INFORMATION

Warranty period is 3 years from the date of purchase for monitor and 1 year for cuff and adapter. This warranty doesn't cover any damages caused by improper using, and also battery, and packaging. When a manufacturing defect is revealed during the warranty period a faulty unit would be repaired or, if repairing is impossible, replaced with another one.

Manufacturing date is in a serial number: WWYYXXXXX.

The manufacturer may change units partially or completely if necessary, without prior notice.

13. ELECTROMAGNETIC COMPATIBILITY INFORMATION

Table 1
For all ME EQUIPMENT and ME SYSTEMS

Guidance and manufacturer's declaration – electromagnetic emissions

<p>The PRO-36 is intended for use in the electromagnetic environment specified below. The customer or the user of the PRO-36 should assure that it is used in such an environment.</p>		
Emissions test	Compliance	Electromagnetic environment-guidance
RF emissions CISPR 11	Group 1	<p>The PRO-36 uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.</p> <p>The PRO-36 is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.</p>
RF emissions CISPR 11	Class B	
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	

Table 2
For all ME EQUIPMENT and ME SYSTEMS

Guidance and manufacturer's declaration – electromagnetic immunity

The PRO-36 is intended for use in the electromagnetic environment specified below. The customer or the user of the PRO-36 should assure that it is used in such an environment.			
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment-guidance
Electrostatic discharge (ESD) IEC 61000-4-2	± 6 kV contact ± 8 kV air	± 6 kV contact ± 8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast transient/ burst IEC 61000-4-4	± 2 kV for power supply lines ± 1 kV for input/output lines	± 2 kV for power supply lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	± 1 kV line(s) to line(s) ± 2 kV line(s) to earth	± 1 kV line(s) to line(s) ± 2 kV line(s) to earth	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<5 % U_T (>95 % dip in U_T) for 0.5 cycle 40 % U_T (60 % dip in U_T) for 5 cycles 70 % U_T (30 % dip in U_T) for 25 cycles <5 % U_T (>95 % dip in U_T) for 5 s	<5 % U_T (>95 % dip in U_T) for 0.5 cycle 40 % U_T (60 % dip in U_T) for 5 cycles 70 % U_T (30 % dip in U_T) for 25 cycles <5 % U_T (>95 % dip in U_T) for 5 s	Mains power quality should be that of a typical commercial or hospital environment. If the user of the PRO-36 requires continued operation during power mains interruptions, it is recommended that the PRO-36 be powered from an uninterruptible power supply or a battery.


Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
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Table 3

For ME EQUIPMENT and ME SYSTEMS that are not LIFE-SUPPORTING

Guidance and manufacturer's declaration – electromagnetic immunity

The PRO-36 is intended for use in the electromagnetic environment specified below. The customer or the user of the PRO-36 should assure that it is used in such an environment.

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment-guidance
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	3 Vrms	Portable and mobile RF communications equipment should be used no closer to any part of the PRO-36, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter Recommended separation distance $d=1.2\sqrt{P}$ $d=1.2\sqrt{P}$ 80 MHz to 800 MHz $d=2.3\sqrt{P}$ 800 MHz to 2,5 GHz <i>where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in metres (m).</i> Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey ^a , should be less than the compliance level in each frequency range ^b . Interference may occur in the vicinity of equipment marked with the following symbol: 
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2,5 GHz	3 V/m	

NOTE 1 At 80 MHz and 800 MHz, the higher frequency range applies.

NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

- a. Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the PRO-36 is used exceeds the applicable RF compliance level above, the PRO-36 should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the PRO-36.
- b. Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.

Table 4
For ME EQUIPMENT and ME SYSTEMS that are not LIFE-SUPPORTING

Recommended separation distances between portable and mobile RF communications equipment and the PRO-36

The PRO-36 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the PRO-36 can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the PRO-36 as recommended below, according to the maximum output power of the communications equipment.

Rated maximum output power of transmitter W	Separation distance according to frequency of transmitter m		
	150 kHz to 80 MHz $d = 1.2 \sqrt{P}$	80 MHz to 800 MHz $d = 1.2 \sqrt{P}$	800 MHz to 2.5 GHz $d = 2.3 \sqrt{P}$
0,01	0,12	0,12	0,23
0,1	0,38	0,38	0,73
1	1,2	1,2	2,3
10	3,8	3,8	7,3
100	12	12	23

*For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (**m**) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in watts (**W**) according to the transmitter manufacturer.*

NOTE 1 At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

14. LATEST REVISION

Latest revision: 2016-02-10

