

SD1 Ultrasonic Pocket Doppler User Manual

About this Manual
P/N: 01.54.458123
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Statement
This manual will help you understand the operation and maintenance of the product better. It is reminded that the product shall be used strictly complying with this manual. User's operation failing to comply with this manual may result in malfunction or accident for which the manufacturer cannot be held liable. The manufacturer owns the copyrights of this manual. Without prior written consent of the manufacturer, any materials contained in this manual shall not be photocopied, reproduced or translated into other languages.

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Responsibility of the Manufacturer
The manufacturer only considers itself responsible for any effect on safety, reliability and performance of the equipment if:
Assembly operations, extensions, re-adjustments, modifications or repairs are carried out by persons authorized by the manufacturer, and
The electrical installation of the relevant room complies with national standards, and
The instrument is used in accordance with the instructions for use.

The manufacturer will make available on request circuit diagrams, component part lists, descriptions, calibration instructions, or other information that will assist service personnel to repair those parts of the equipment that are designated by the manufacturer as repairable by service personnel.

Product Information
Product Name: Ultrasonic Pocket Doppler
Model:SD1

Terms Used in this Manual
This guide is designed to give key concepts on safety precautions.

WARNING
WARNING label advises against certain actions or situations that could result in personal injury or death.

CAUTION
A **CAUTION** label advises against actions or situations that could damage equipment, produce inaccurate data, or invalidate a procedure.

NOTE
A **NOTE** provides useful information regarding a function or a procedure.

● **Safety Precautions**

CAUTION
Federal (U.S.) Law restricts this device to sale by or on the order of a physician.

NOTE:
This user manual is written to cover the maximum configuration. Therefore, your model may or may not have some of the parameters and functions described, depending on what you have ordered.

This unit is internally powered equipment, and it is an IEC/EN 60601-1 Type BF applied part. Type BF protection means that the connection between the equipment and personnel complies with permitted leakage currents and dielectric strength of IEC/EN 60601-1.

WARNING and **CAUTION** messages must be observed. To avoid the possibility of injury, observe the following precautions during the operation of the device.

WARNING
1 It is to be used by health care professionals on the order of a physician.

2 The Doppler is a tool to aid the healthcare professional in hospitals, clinics and should not be used in place of normal fetal detection. It is not intended for treatment or use during labor and delivery.

3 Placement of the ultrasound transducer on the abdomen is critical to obtaining the fetal heart beat as opposed to maternal heart beat or other abdominal noise. The user should be trained in proper placement techniques either through acceptable Ob/Gyn training and individual state accreditation, or as being prescribed by such a trained clinician and trained in device placement.

4 This device is not explosion-proof and cannot be used in the presence of flammable anesthetics.

5 Magnetic and electrical fields are capable of interfering with the proper performance of the device. For this reason, make sure that all external devices operated in the vicinity of this device comply with the relevant EMC requirements. X-ray equipment and magnetic resonance imaging (MRI) devices can emit high levels of electromagnetic radiation.

6 We recommend that exposure to ultrasound should be kept as low as reasonably achievable. This is considered to be good practice and should be observed at all time.

7 Do not use the device with HF surgical equipment and do not use it in an MRI environment.

8 The device is not protected against defibrillation.

9 **SHOCK HAZARD - Do not attempt to replace batteries with wet hands.**

10 The device should not be used adjacent to or stacked with other equipment and that if adjacent or stacked use is necessary, the device should be observed to verify normal operation in the configuration in which it will be used.

11 The medical electrical equipment needs to be installed and put into service according to the EMC Information provided in this user manual.

12 Portable and mobile RF communications equipment can affect medical electrical equipment; refer to section Recommended Separation Distances.

13 Do not service or maintain the device or any accessory which is in use with a patient.

CAUTION

1 Refer servicing to qualified personnel.

2 Keep the device in a clean environment and avoid vibration during storage.

3 Do not sterilize the Doppler.

4 Electromagnetic Interference - Ensure that the environment in which the device is operated is not subject to any source of strong electromagnetic emissions, such as radio transmitters, mobile telephones, etc.

5 Prior to examination using the Doppler, check for visible damages of the main unit and the probe that may endanger the patient/operator or machine performance. If the damage is found, replace them with good ones at once.

6 The following safety checks should be performed once every two years or as specified in the institution's test and inspection protocol by a qualified person who has adequate training, knowledge, and practical experience to perform these tests.

◆ Inspect the equipment for mechanical and functional damage.
◆ Inspect the safety relevant labels for legibility.
◆ Inspect the equipment for mechanical and functional damage.
◆ Inspect the safety relevant labels for legibility.

The leakage current should never exceed the limit. The data should be recorded in an equipment log. If the device is not functioning properly or fails any of the above tests, the device has to be repaired.

7 The device and accessories are to be disposed of according to local regulations after their useful lives. Alternatively, they can be returned to the dealer or the manufacturer for recycling or proper disposal. Batteries are hazardous waste. Do NOT dispose them together with house-hold garbage.

● **Introduction**

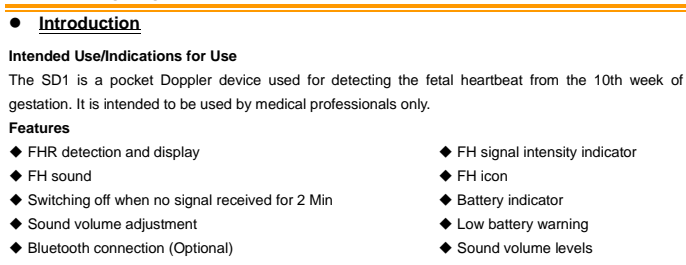
Intended Use/Indications for Use
The SD1 is a pocket Doppler device used for detecting the fetal heartbeat from the 10th week of gestation. It is intended to be used by medical professionals only.

Features

◆ FHR detection and display
◆ FH sound
◆ Switching off when no signal received for 2 Min
◆ Sound volume adjustment
◆ Bluetooth connection (Optional)

◆ FH signal intensity indicator
◆ FH icon
◆ Battery indicator
◆ Low battery warning
◆ Sound volume levels

Appearance (Above pictures are just for reference)



LCD Display & Touch Keys



| Item | Description |
|------|--|
| 1 | Fetal heart icon Indicates fetal heart beat and flickers to the fetal heart beat. |
| 2 | Fetal heart signal intensity indicator This indicator displays on the left side of the screen and has three status: empty, half empty and full, which respectively represents low, medium and high fetal heart signal intensity. |
| 3 | FHR numeric Displays fetal heart rate within the range from 50 bpm to 240 bpm. When fetal heart rate is out of the range, it displays "--". Sound volume numeric Sound volume numeric is displayed in the center of the screen, the same area as the FHR numeric. When you adjust sound volume, the sound volume numeric will display for 0.5 second before switching back to display FHR numeric. Sound volume ranges from level 0 to 7. |
| 4 | Battery indicator Battery indicator displays on the right side of the screen. There are 5 battery levels, represented by 0 to 4 panes in the icon. When battery is empty, battery empty icon will be displayed and flickering, and the battery needs replacing. |
| 5 | Sound volume increase touch key Touch the key to increase sound volume. |
| 6 | Sound volume decrease touch key Touch the key to decrease sound volume. |
| 7 | On/Off touch key When the Doppler is off, touch this key for a little while to switch it on; When the Doppler is on, touch this key for a little while to switch it off. |

Battery
SD1 is powered by two AA alkaline batteries. Battery specification: LR6, AA, 1.5 V;

Note:
You can use AA alkaline batteries of the same specification purchased locally.

● **Basic Operation**

NOTE:
To ensure that the Doppler works properly, please read this chapter and Chapter Safety Precautions before operation; follow the steps when connecting all the components.

Opening the Package and Checking
Open the package; take out the Doppler and accessories carefully. Keep the package for possible future transportation or storage. Check the components according to the packing list.

◆ Check for any mechanical damage.
◆ Check all the cables and accessories.
If there is any problem, contact us or your local distributor immediately.

Installing the Battery
a) Unscrew the screw with a cross screwdriver and remove the battery compartment cover.
b) Insert the battery into the compartment carefully. Ensure its anode and cathode terminals are aligned with the anode and cathode marks on the compartment.
c) Install the compartment cover and secure it with the screw.

Removing/Replacing the Battery
a) Unscrew the screw with a cross screwdriver and remove the battery compartment cover.
b) Take out the used battery. You can also replace it with a new one. Ensure the new battery's terminals are placed in the right direction as indicated by the anode and cathode marks.
c) Install the compartment cover and secure it with the screw.

WARNING

1 Turn off the Doppler before removing or replacing the battery.
2 Replace alkaline batteries with those of identical specifications provided by the manufacturer or purchased locally. See Chapter Product Specifications for details about battery specifications.

3 If the batteries have been inserted incorrectly, the Doppler will not function or it will be damaged.

4 Do not disassemble or short-circuit batteries.
5 Do not recharge batteries.
6 Do not dispose of batteries in fire or water.
7 Do not allow metal objects to contact the battery terminals.
8 Do not mix with used or other battery type (such as alkaline with carbon zinc).
9 Do not solder the batteries directly. If soldering or welding connection to the battery is required, consult our engineer for proper methods.
10 Do not over-discharge batteries.
11 To install or remove batteries, follow the equipment manufacturer's instructions.
12 Keep battery away from small children. If swallowed, consult a physician at once.
13 Store the battery in cool, dry place before use. Do not keep batteries at temperature of 45°C or above, or at humidity of 75% or above.
14 Dispose the battery according to the local regulations. Refer to IEC61429 for standard disposal when necessary.
15 Remove the battery and store it at a cool and dry environment if the Doppler is not used for a long time.
16 Batteries have life cycles. If the time that the Doppler using battery becomes much shorter than usual, the battery life is at an end. Replace the battery with a new one of the same specification as the one provided or recommended by the manufacturer.

Switching On
Touch the On/Off touch key for about 1second when the Doppler is off, and the Doppler will display the switching on interface before switching to display the test interface.

Switching Off
Touch the On/Off touch key for about 1second when the Doppler is on, and the Doppler will be switched off.
If the Doppler is not in operation or no signal is received for 2 minutes, the Doppler will switch off automatically.

● **FHR Detection**

Before applying the Doppler to inspect FHR, you should always check whether the Doppler is in good condition and whether there is evident damage that might affect patient's safety and the device's function. If evident damage is found, stop using it at once and replace it with a good one.

Procedures to Detect FHR:

a) Have the patient lie on her back.
b) Apply appropriate amount of coupling gel to the ultrasonic transducer head of the Doppler and switch on the Doppler.
c) Palpate the patient's abdomen gently to confirm the fetus's position.
d) Place the Doppler on the patient's abdomen, and move it around the fetus's position or tilt it until a clear and rhythmic heart sound is heard and FHR numeric is stably displayed.

Note:
1 Do not mistake the maternal heart rate for fetal heart rate. The fetal pulse should be different from the maternal pulse, which can be measured at the wrist or neck.
2 Do not wear gloves to touch the keys. If there's water and coupling gel on the fingers, please clean them first or the touching effect will be influenced.

How to Find the Best FH Signal:
1) The easiest way: take the position the doctor last detected for FHR as a reference and move the Doppler around the position slowly until the best FH signal is found.
2) The fetal heart position may change as the fetus moves inside the uterus. You can confirm the fetal position first according to the position of the uterus fundus in different gestational weeks.
The clearest and loudest fetal heart sound is generally obtained when the Doppler is placed on the fetus's back. Fetal movement is usually the movement of fetal limbs. So, if frequent fetal movement occurs at the right side of the abdomen, the fetus's back is probably at the left side and vice versa. You can find the fetus's back according to fetal movement's position.
If the fetus is in cephalic delivery position, the fetal heart is either on the right side or on the left side below the navel; if the fetus is in breech delivery position, the fetal heart is either on the right side or on the left side above the navel.

Steps to Find Fetal Heart:
Have the patient lie on back and relax >> confirm fetal position by hand >> apply coupling gel to the Doppler >> place the Doppler on patient's abdomen and start looking for the fetal heart >> the fetal heart is found when the Doppler gives out a continuing thumping sound "boom-boom-boom".

CAUTION

1 The Doppler's degree of protection against harmful ingress of water is IP22. Do not immerse it in water.
2 The Doppler is delicate and sensitive. Please handle it with care and try to avoid dropping on to the ground or any hard surfaces. Any damage caused by dropping is not covered by the warranty.
3 Keep the coupling gel away from children. If swallowed, consult a physician at once.

Note:
1 The best quality of fetal heart signal is obtained only when the Doppler is placed on the best detection position.
2 Do not place the Doppler near positions where placental sound or umbilical blood flow sound is loud.
3 If the fetus is in the cephalic position and the mother is supine, the clearest heart sound will normally be found on the midline below the navel. During detection, the pregnant woman's prolonged lying in the supine position should be avoided to reduce the possibility of supine hypotension. Putting a pillow or cushion under the patient's head or feet can be of help.
4 It is not possible to obtain accurate FHR unless a clear fetal heart signal is detected. If the calculated FHR is not in accordance with the beat of the fetal heart sound, the fetal heart sound auscultation result shall prevail.
5 When applied to the patient, the Doppler may warm slightly (less than 2°C (35.6°F) above ambient temperature). When NOT applied, the Doppler may slightly (less than 5°C (41°F) above ambient temperature).

After Detection
1) Switch off the Doppler.
2) Wipe the remaining gel off the patient and the probe with a clean soft cloth or tissue.

● Mobile Application Software (APP)

SD1 can connect to mobile phones with its Bluetooth function (optional). The SD1 Care APP has both Android and iOS versions.

iOS APP operating environment:
A) hardware environment
Processor: dual-core Apple A6
RAM: ≥1GB
B) software environment: iOS 8.0 and above operating system
C)network environment: support Bluetooth

Android APP operating environment:
A) hardware environment
CPU: frequency≥1.0GHz
RAM: ≥1GB
B)software environment: Android 4.3 and above operating system
C)network environment: support Bluetooth

How to use SD1 Care APP
1. Download and install software
Scan either of the following QR codes to download the SD1Care APP, and install and run it as prompted.

Note:
1 Your mobile phone may prohibit the installation of "applications from unknown sources". Enter **Settings** to allow the installation first.
2 For normal functioning of the APP, please give the APP function-related permissions.
3 For how to use the APP, read the instructions in the **About** sub-interface under the **Settings** interface of the APP.

2. Activate the device
Open the APP and go to Settings>Activation and input SD1 activation code (14 numbers after 01).

3. Pair device
Open Bluetooth function of the mobile to automatically pair the SD1.

4. Start detection
Put the coupling gel on SD1 and position the probe to the optimal place of maternity's abdomen. And click the "start" key. After pressing start, confirm that the data on the APP and the SD1 probe match. As with any Bluetooth communication, it is important to make sure the connection is not compromised.

5. Adjust the fetal heart beat sound volume
When using mobile phone to play the fetal heart beat sound, you can adjust the volume with the volume keys of the mobile phone. When using SD1 to play the heart beat sound, touch 'volume+' or 'volume-' to adjust the volume.

6. Finish the detection
When the detection is finished, click 'Stop' touch key and the detection data will be saved automatically. Note: Please make sure your mobile phone has enough battery power, and avoid killing the process directly or switching to other applications during the fetal heart detection.

7. Real time detection mode and DEMO mode
We provide DEMO mode for users' reference. You can turn on DEMO key in Setup and enter fetal heart detection interface to watch the DEMO. The word 'DEMO' is displayed in the interface to distinguish from real time detection.

WARNING
SD1 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.

NOTE:
1. This equipment (SD1) 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.
2. Any changes or modifications to this unit not expressly approved by the party responsible for compliance could void the user's authority to operate the equipment.

● **Maintenance and Cleaning**

Maintenance
Before each use, check if the equipment has visible evidence of damage that may affect the patient and the operator's safety or the Doppler's functioning. If the damage is evident, contact the manufacturer for service or replace it.

The overall check of the Doppler, including safety check and function check, should be performed by qualified personnel every 12 months, and each time after service. And safety check must include current leakage test and insulation test. Besides the above requirements, comply with local regulations on maintenance and measurement.

The accuracy of FHR is determined by the Doppler and cannot be adjusted by user. If you have doubt concerning the accuracy of FHR, verify it with other methods such as using a stethoscope, or contact local distributor or the manufacturer for help.

The Doppler is fragile and must be handled with care. Wipe the remaining gel off the Doppler after each use. These measures can help prolong the Doppler's life.

Replace the accessories such as the battery according to use. If any of the accessories are damaged, refer to chapter **Ordering Information** for details and order new ones.
Please check the label for the date of manufacture, the service life is 5 years (The service life is limited to the Doppler, not including the replaceable accessories. The only replaceable accessory of SD1 is battery. The frequency of usage is 8 hours/day).

Cleaning
Before cleaning, switch off the Doppler.
Keep the exterior surface of the device clean and free of dust and dirt.
Clean the exterior surface of the Doppler with a dry, soft cloth. If necessary, clean it using a soft cloth dampened with mild neutral detergent, ethanol (75%) or isopropanol (70%), and then wipe it dry with a dry cloth immediately.

CAUTION

1 Do not use strong solvent, such as acetone.
2 Never use an abrasive such as steel wool or metal polish.
3 The Doppler's degree of protection against harmful ingress of water is IP22. Do not immerse it in water.
4 Do not remain any solution on the surface after cleaning.

Disinfection
In normal use the Doppler does not need disinfection. In case of being soiled, clean the main unit case and then disinfect it by wiping it with a soft cloth dampened with ethanol (75%) or isopropanol (70%). Then wipe it dry with a dry cloth.

CAUTION
Do not immerse the Doppler into the disinfectant.

● **Product Specifications**

Product Information

Product Name Ultrasonic Pocket Doppler
Model SD1

Complied Standards
IEC 60601-1:2005/A1:2012, EN 60601-1:2006/A1:2013, IEC 60601-1-2:2014, IEC 60601-2-37:2015, IEC 61266:1994

Classification
Anti-electric Shock Type: Internally powered equipment
Anti-electric Shock Degree: Type BF equipment

Degree of Protection against Harmful Ingress of Water: IP22. Protection against vertically falling water drops when ENCLOSURE tilted up to 15°

Degree of Safety in Presence of Flammable Gases: Equipment not suitable for use in presence of flammable gases

Working System: Continuous running equipment
EMC: CISPR 11 Group 1 Class B

Physical Specifications

Size: Length*Width* Height: (48±2) mmx (39±2) mmx (147±3) mm
Weight: < 180g

LCD: Size: (24±2) mmx (13±2) mm
Display: ◆FHR
◆Battery level
◆Signal intensity
◆Sound volume level
◆FH icon

Coupling Gel: pH: 5.5-8.0
Acoustic Impedance: 1.5x10⁶ Pa.s/m ~1.7x10⁶ Pa.s/m (35°C/95°F)

Environment

Working: Temperature: +5°C ~ +40°C (+41°F ~ +104°F)
Humidity: 15% RH ~ 95% RH (non-condensing)
Atmospheric Pressure: 70kPa ~ 106 kPa

Transport and Storage: Temperature: -25°C ~ +70°C (-13°F ~ +158°F)
Humidity: 15% RH ~ 95% RH (non-condensing)
Atmospheric Pressure: 70 kPa ~ 106 kPa

Note: The time required for the Doppler to warm from the minimum storage temperature between uses until it is ready for intended use is at least 2 hours; the time required for the Doppler to cool from the maximum storage temperature between uses until it is ready for intended use is at least 2 hours

Performance Specifications

FHR Performance: (Essential) FHR Measuring Range: 50 bpm ~ 240 bpm
Accuracy: ±2 bpm
Note: FHR measurement result may not be accurate if the equipment is measuring beyond its measuring range.

FHR Resolution: 1bpm

Audio Output: Output Power: 2w
Background noise: <45dBA

Auto Power-off: Power off when the Doppler receives no signal or operation for 2 minutes.

Bluetooth: Transmission Range (Without Obstacles) :>5m (Indoor range depends on the building's structure and material.)
Nominal Frequency: 3MHz
Working Frequency: 3MHz

Ultrasound: p < 1 MPa
Iob<10 mW/cm2
Isp1a<100 mW/cm2
Isata<10 mW/cm2
Isp3a<190 W/cm2
Isp1a.3<94 mW/cm2
Effective Radiating Area: 490mm2 ± 15%
Working Mode: pulse wave

Battery Specifications

| | | | | | |
|--|--|---------|----------|------|---------------------------------------|
| Specification: | Two AA 1.5V alkaline batteries (AA, LR6, 1.5V) | | | | |
| Working Duration: | ◆ ≥26h | | | | |
| Bluetooth Specifications | | | | | |
| FCC ID | SMQSD1 | | | | |
| Modulation: | GFSK | | | | |
| Frequency: | 2400-2483.5MHz | | | | |
| Tolerance Frequency: | ± 20ppm | | | | |
| RF output power: | ≤ 200mW (EIRP) | | | | |
| Occupied Channel Bandwidth: | ≤ 3MHz | | | | |
| Transmitter Unwanted Emissions: | ≤ -30dBm | | | | |
| Low Output Summary Table (For systems whose global maximum value does not exceed 1.0) System: SD1 Ultrasonic Pocket Doppler | | | | | |
| Model (MHz) | $I_{p_{max}}$ (mW/cm ²) | TI Type | TI Value | MI | $I_{p_{max}}$ (W/cm ²) |
| SD1 CD3.0 | 5.69 | TIS | 0.05 | 0.01 | 0.02 |
| | | TIB | 0.01 | | |

● **Ordering Information**

CAUTION
Only the parts supplied by the manufacturer should be used with the Doppler.

| Parts | Part Number |
|------------------------------------|--------------|
| Main Unit | |
| SD1 Doppler(Non-Bluetooth version) | 02.06.262535 |
| SD1 Doppler(Bluetooth version) | 02.06.262639 |
| Accessories | |
| AA Alkaline Battery | 01.21.064086 |
| Normal Carry Case | 01.56.466428 |
| Coupling Gel | 01.57.078170 |

● **Ultrasound Intensity and Safety**

Ultrasound in Medicine
The use of diagnostic ultrasound has proved to be a valuable tool in medical practice. Given its known benefits for non-invasive investigations and medical diagnosis, including investigation of the human fetus, the question of clinical safety with regards to ultrasound intensity arises. There is no easy answer to the question of safety surrounding the use of diagnostic ultrasound equipment. Application of the ALARA (As Low As Reasonably Achievable) principle serves as a rule-of-thumb that will help you to get reasonable results with the lowest possible ultrasound output. The American Institute of Ultrasound in Medicine (AIUM) states that given its track record of over 25 years of use and no confirmed biological effects on patients or instrument operators, the benefits of the prudent use of diagnostic ultrasound clearly outweigh any risks.

Ultrasound Safety and the ALARA Principle
Ultrasound waves dissipate energy in the form of heat and can therefore cause tissue warming. Although this effect is extremely low with Doppler, it is important to know how to control and limit patient exposure. Major warming is related to the type of tissue. Although no confirmed adverse mechanical effects on patients or mammals caused by exposure at intensities typical of present diagnostic ultrasound instruments have ever been reported, the threshold for cavitation is still undetermined. Generally speaking, the higher the acoustic pressure, the greater the potential for mechanical bioeffects; the lower the acoustic frequency, the greater the potential for mechanical bioeffects.

The AIUM and NEMA formulate mechanical index (MI) in order to indicate the potential for mechanical effects. The MI is defined as the ratio of the peak-rarefactional acoustic pressure (should be calculated by tissue acoustic attenuation coefficient 0.3 dB/cm/MHz) to the acoustic frequency.
 $MI = \frac{Pr}{fawt} \times CMI$
 $CMI = 1 \text{ (MPa / MHz)}$

TI (Thermal Index)
Heating of tissues is caused by absorption of ultrasound when the ultrasound energy is applied. The temperature rise is determined by the acoustic intensity, exposed area and thermo physical properties of the tissue.

In order to indicate the potential for temperature rise caused by thermal effects, the AIUM and NEMA formulate thermal index (TI) which is defined as the ratio of the total acoustic power to the acoustic power required to raise the tissue temperature by 1°C (1.8°F). According to different thermo physical properties of the tissue, TI is divided into three kinds: TIS, TIB and TIC.
TIS (Soft Tissue Thermal Index): It provides an estimate of potential temperature rise in soft or similar tissues.
TIB (Bone Thermal Index): It provides an estimate of potential temperature rise when the ultrasound beam passes through soft tissue and a focal region is in the immediate vicinity of bone.
TIC (Cranial Bone Thermal Index): It provides an estimate of potential temperature rise in the cranial bones or superficial bones.

Measurement Uncertainties
The uncertainties in the measurements were predominantly systematic in origin; the random uncertainties were negligible in comparison. The overall systematic uncertainties were determined as follows:

1. Hydrophone Sensitivity: ± 12percent for intensity, ± 6 percent for pressure. Based on the hydrophone calibration report by ONDA. The uncertainty was determined within ± 1 dB in frequency range 1-15 MHz.
2. Digitizer: ± 0.3 percent for intensity, ± 0.15 percent for pressure.
3. Based on the stated accuracy of the 8-bit resolution of the Agilent DSO6012 Digital Oscilloscope and the signal-to-noise ratio of the measurement.
4. Temperature: ± 2.4 percent for intensity uncertainty, ± 1.2 percent for pressure uncertainty. Based on the temperature variation of the water bath of ± 1°C (1.8°F).
5. Spatial Averaging: ± 3.5 percent for intensity, ± 1.75percent for pressure.
6. Non-linear Distortion: N/A.
7. No effects of nonlinear propagation were observed.

Since all the above error sources are independent, they may be added on an RMS basis, giving a total uncertainty of ± 12.73 percent for all intensity values reported, ± 6.37 percent for all the pressure values, ± 12.6 percent for the Mechanical Index, uncertainty of ± 12.73% percent for power, ± 0.15 percent for center frequency, ± 0.87% for the MI.

Prudent Use Statement
Although no confirmed bioeffects on patients caused by exposure from present diagnostic ultrasound equipment have ever been reported, the potential exists that such bioeffects may be identified in the future. Therefore, the ultrasound should be used prudently. High levels of acoustic output and long exposure time should be avoided while acquiring necessary clinical information.
Reference for Acoustic Output and Safety
1. "Bioeffects and Safety of Diagnostic Ultrasound" issued by AIUM in 1993
2. "Medical Ultrasound Safety" issued by AIUM in 1994
3. "Acoustic Output Measurement Standard for Diagnostic Ultrasound Equipment, Revision 3" issued by AIUM/NEMA in 2004
4. "Standard for real-time display of thermal and mechanical acoustic output indices on diagnostic ultrasound equipment, Revision 2" issued by AIUM/NEMA in 2004
5. "Information for Manufacturers Seeking Marketing Clearance of Diagnostic Ultrasound Systems and Transducers" issued in 2008.
6. "Medical electrical equipment—Part 2-37: Particular requirements for the basic safety and essential performance of ultrasonic medical diagnostic and detection equipment" issued by IEC in 2007.

Acoustic Output Reporting Table for Track 1 Acoustic output reporting table for IEC60601-2-37(IEC60601-2-37, Edition 2.1, 2015-0, table 201.103)

| Index label | MI | TIS | | TIB | | TIC |
|------------------------------|--|------------|---------------|------------|---------------|-----|
| | | At Surface | Below Surface | At surface | Below Surface | |
| Maximum index value | 0.01 | 0.05 | | 0.01 | | N/A |
| Index component value | | N/A | 0.05 | NA | 0.01 | |
| Acoustic Parameters | Pr_{at} (MPa) | 0.02 | | | | |
| | P (mW) | | 7.35 | | 7.35 | N/A |
| | P_{T1} (mW) | | N/A | | N/A | |
| | zS (cm) | | | 3.50 | | |
| | zB (cm) | | | | 3.70 | |
| | zMI (cm) | 3.70 | | | | |
| | zP_{II} (cm) | 3.70 | | | | |
| | $fawf$ (MHz) | 3.00 | 3.00 | | 3.00 | N/A |
| | pr (Hz) | 5000 | | | | |
| | sr (Hz) | N/A | | | | |
| Other information | $npds$ | 1 | | | | |
| | lpa at zP_{II} (W/cm ²) | 0.02 | | | | |
| | $Ispta$ at zP_{II} or $zSII$ (mW/cm ²) | 5.69 | | | | |
| | $Ispta$ at zP_{II} or $zSII$ (mW/cm ²) | 12.26 | | | | |
| | pr at zP_{II} (MPa) | 0.04 | | | | |
| Operating control conditions | Fixed | | | | | |

Acoustic Output Reporting Table for Track1(Non-autoscanning Mode)

| Acoustic Output | | | | | |
|-------------------------------|-----------------|-------------------------------|------------------------------|-------|--|
| Global Maximum Value | MI | ISPTA.3 (mW/cm ²) | ISPPA.3 (W/cm ²) | | |
| Pr_3 (MPa) | 0.02 | 0.01 | 5.69 | 0.02 | |
| W_0 (mW) | | 7.35 | 8.97 | | |
| I_c (MHz) | 3.00 | 3.00 | 3.00 | | |
| Zsp (cm) | 3.70 | 3.70 | 3.70 | | |
| Associated Acoustic Parameter | Beam dimensions | X-6 (cm) | 2.50 | 2.50 | |
| | | Y-6 (cm) | 2.50 | 2.50 | |
| | PD (usec) | 72.25 | 72.25 | 72.25 | |
| | PRF (Hz) | 5000 | 5000 | 5000 | |
| | EBD (cm) | | 2.50 | | |
| Operating Control Conditions | Fixed | | | | |
| | | | | | |

Standard Parameter Equal Contrast List IEC60601-2-37 Standard Parameters

| Parameter | Note | Parameter | Note |
|---------------|---|----------------------|--|
| P_{ra} | Attenuated Peak-rarefactional Acoustic Pressure | f_{awf} | Center Frequency, Acoustic Working Frequency |
| P_r | Peak-rarefactional Acoustic Pressure | X | -12dB Output Beam Dimensions |
| P | Output Power | Y | |
| Z_s | Depth of Soft Tissue Thermal Index | t_d | Pulse Duration |
| $P_o(Z_s)$ | Attenuated Output Power | prf | Pulse Repetition Frequency (Pulse Repetition Rate) |
| $I_{ms}(Z_s)$ | Attenuated Temporal-average Intensity | d_{eq} | Equivalent Beam Diameter |
| Z_{sp} | Break-point Depth | $l_{p,at}$ at max MI | Attenuated Intensity at the point of Maximum MI |
| Z_b | Depth for Bone Thermal Index | A_{apert} | -12dB Output Beam Area |
| $I_{p,at}$ | Attenuated Pulse-intensity Integral | MI | Mechanical Index |
| $I_{p,at}$ | Attenuated Intensity Integral | TIS | Soft Tissue Thermal Index |
| $d_{eq}(Z_s)$ | Equivalent Beam Diameter at the point of Z_{sp} | TIB | Bone Thermal Index |
| | | TIC | Cranial-bone Thermal Index |

● **EMC Information**

Electromagnetic Emissions
Guidance and manufacturer's declaration – electromagnetic emission
The SD1 Ultrasonic Pocket Doppler is intended for use in the electromagnetic environment specified below. The customer or the user of the device should assure that it is used in such an environment.

| RF emissions | Compliance | Electromagnetic environment - guidance |
|--------------|------------|--|
| CISPR 11 | Group 1 | The SD1 Ultrasonic Pocket Doppler 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. |

| RF emission | Class | Guidance and manufacturer's declaration – electromagnetic immunity |
|--|----------------|--|
| CISPR 11 | Class B | The SD1 Ultrasonic Pocket Doppler 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. |
| Harmonic emissions IEC/EN61000-3-2 | Not applicable | |
| Voltage fluctuations / flicker emissions IEC/EN61000-3-3 | Not applicable | |

Electromagnetic Immunity

Guidance and manufacturer's declaration – electromagnetic immunity
The SD1 Ultrasonic Pocket Doppler is intended for use in the electromagnetic environment specified below. The customer or the user of the device 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 | ±8kV contact ±15kV air | ±8kV contact ±15kV air | Floors should be wood, concrete or ceramic tile. If floor are covered with synthetic material, the relative humidity should be at least 30%. |
| Electrical Fast Transient/Burst IEC/EN61000-4-4 | ±2kV for power supply lines ±1kV for input/output lines | Not applicable | Not applicable |
| Surge IEC/EN61000-4-5 | ±1 kV line(s) to line(s) ± 2 kV line(s) to earth | Not applicable | Not applicable |
| Voltage dips, short interruptions, and voltage variations on power supply input lines IEC/EN61000-4-11 | <5%UT(>95% dip in UT) for 0.5cycle 40%UT(60% dip in UT) for 5 cycles 70%UT(30% dip in UT) for 25 cycles <5%UT(>95% dip in UT) for 5s | Not applicable | Not applicable |
| Power frequency (50Hz/60Hz) magnetic field IEC61000-4-8 | 30 A/m | 30 A/m | Power magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment. |
| IMMUNITY to proximity magnetic fields | 8A/m, Modulation: CW Test frequency:30KHz; 65A/m, Modulation: Pulse modulation, 2.1KHz Test frequency:134.2KHz; 7.5A/m, Modulation: Pulse modulation, 50KHz Test frequency:13.56MHz; | 8A/m, Modulation: CW Test frequency:30KHz; 65A/m, Modulation: Pulse modulation, 2.1KHz Test frequency:134.2KHz; 7.5A/m, Modulation: Pulse modulation, 50KHz Test frequency:13.56MHz; | |

Electromagnetic Immunity

Guidance and manufacturer's declaration – electromagnetic immunity
The SD1 Ultrasonic Pocket Doppler is intended for use in the electromagnetic environment specified below. The customer or the user of the device should assure that it is used in such an environment.

| Immunity test | IEC 60601 test level | Compliance level | Electromagnetic environment-guidance |
|---------------------------|--|-----------------------------|--|
| Conducted RF IEC61000-4-6 | 3 V _{rms} 150 kHz ~ 80 MHz 6V(rms) in ISM bands between 0,15 MHz and 80 MHz | Not Applicable | Portable and mobile RF communications equipment should be used no closer to any part of the SD1 Ultrasonic Pocket Doppler, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance: $d = 0.35\sqrt{P}$ 80 MHz to 800 MHz $d = 0.7\sqrt{P}$ 800 MHz to 2.7 GHz $d = 6\sqrt{P} / E$ at RF wireless communications equipment bands (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 SD1 Ultrasonic Pocket Doppler, including cables specified by the manufacturer). 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 meters (m). 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: (img) |
| Radiated RF IEC61000-4-3 | 10V/m 80 MHz ~ 2.7 GHz | 10 V/m 80 MHz to 2.7 GHz | |

NOTE1: At 80 MHz and 800 MHz, the higher frequency range applies.

NOTE2: 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 SD1 Ultrasonic Pocket Doppler is used exceeds the applicable RF compliance level above, the SD1 Ultrasonic Pocket Doppler should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting or relocating the SD1 Ultrasonic Pocket Doppler.
- b) Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.
- c) The ISM (industrial, scientific and medical) bands between 0,15 MHz and 80 MHz are 6,765 MHz to 6,795 MHz; 13,553 MHz to 13,567 MHz; 26,957 MHz to 27,283 MHz; and 40,66 MHz to 40,70 MHz. The amateur radio bands between 0,15 MHz and 80 MHz are 1,8 MHz to 2,0 MHz, 3,5 MHz to 4,0 MHz, 5,3 MHz to 5,4 MHz, 7 MHz to 7,3 MHz, 10,1 MHz to 10,15 MHz, 14 MHz to 14,2 MHz, 18,07 MHz to 18,17 MHz, 21,0 MHz to 21,4 MHz, 24,89 MHz to 24,99 MHz, 28,0 MHz to 29,7 MHz and 50,0 MHz to 54,0 MHz.

Table-Test specifications for ENCLOSURE PORT IMMUNITY to RF wireless communications equipment

| Test Frequency (MHz) | Brand ^{a)} (MHz) | Service ^{a)} | Modulation ^{b)} | Maximum Power(W) | Distance (m) | IMMUNITY TEST LEVEL (V/m) |
|----------------------|---------------------------|---|---|------------------|--------------|---------------------------|
| 385 | 380-390 | TETRA 400 | Pulse modulation ^{b)} 18Hz | 1.8 | 0.3 | 27 |
| 450 | 430-470 | GMR5 460, FRS 460 | FM ^{c)} ±5 kHz deviation 1kHz sine | 2 | 0.3 | 28 |
| 710 | 704-787 | LTE Brand 13, 17 | Pulse modulation ^{b)} 217 Hz | 0.2 | 0.3 | 9 |
| 810 | | GSM 800/900, TETR A 800, IDEN 820, CDMA 850, LTE Band 5 | Pulse modulation ^{b)} 18 Hz | 2 | 0.3 | 28 |
| 930 | 800-960 | | | | | |
| 1720 | 1700-1990 | GSM 1800; CDMA 1900; GSM 1900; DECT; LTE Band 1, 3, 4, 25; UMTS | Pulse modulation ^{b)} 217 Hz | 2 | 0.3 | 28 |
| 1845 | | | | | | |
| 1970 | 2400-2570 | Bluetooth, WLAN, 802.11 b/g/n, RFID 2450, LTE Band 7 | Pulse modulation ^{b)} 217 Hz | 2 | 0.3 | 28 |
| 5240 | 5100-5800 | WLAN 802.11 a/n | Pulse modulation ^{b)} 217 Hz | 0.2 | 0.3 | 9 |
| 5500 | | | | | | |
| 5785 | | | | | | |

Note: 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 1m. 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.
- c) As an alternative FM modulation, 50% pulse modulation at 18 Hz may be used because while it does not represent actual modulation, it would be worst case

Recommended Separation Distances

Recommended separation distances between portable and mobile RF communications equipment and the SD1 Ultrasonic Pocket Doppler

The SD1 Ultrasonic Pocket Doppler is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the SD1 Ultrasonic Pocket Doppler can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the SD1 Ultrasonic Pocket Doppler 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 $d = 0.35\sqrt{P}$ | 800 MHz to 2.7GHz $d = 0.7\sqrt{P}$ |
| 0.01 | / | 0.035 | 0.07 |
| 0.1 | / | 0.11 | 0.22 |
| 1 | / | 0.35 | 0.7 |
| 10 | / | 1.11 | 2.21 |
| 100 | / | 3.5 | 7 |

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.

Table 11 – Test specifications for ENCLOSURE PORT IMMUNITY to proximity magnetic fields

| Test frequency | Modulation | IMMUNITY TEST LEVEL (A/m) |
|----------------------|--------------------------------|---------------------------|
| 30 kHz ^{a)} | CW | 8 |
| 134.2 kHz | Pulse modulation ^{b)} | 65 ^{c)} |
| 13.56 MHz | Pulse modulation ^{b)} | 7.5 ^{d)} |

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

● **Overall Sensitivity**

| D | d | A | B | | | V _s | V _n | C | S | |
|------------------------|------|----------------|-----------------|----------------|------|---------------------------|----------------|------|--------|--------|
| | | | γ _{Ba} | B _v | B | | | | | |
| 1.58 3MHz | 50 | 40.9 | T | 5#4# | 0 | 77.0 | 140 | 75 | 5.42 | 123.32 |
| | | | B _a | 77.0 | | | | | | |
| | 75 | 44.4 | T | 5#3# | 0 | 68.4 | 80 | 40 | 6.02 | 118.82 |
| | | | B _a | 68.4 | | | | | | |
| | 100 | 46.9 | T | 5#3# | 0 | 69.6 | 180 | 90 | 6.02 | 122.52 |
| | | | B _a | 69.6 | | | | | | |
| 200 | 52.9 | T | 5#3# | 0 | 68.4 | 83 | 42 | 5.92 | 127.22 | |
| | | B _a | 75.1 | | | | | | | |
| 2.38 3MHz | 50 | 39.0 | T | 5#4# | 0 | 77.0 | 130 | 69 | 5.50 | 121.50 |
| | | | B _a | 77.0 | | | | | | |
| | 75 | 42.5 | T | 5#4# | 0 | 77.0 | 115 | 55 | 6.41 | 125.91 |
| | | | B _a | 77.0 | | | | | | |
| | 100 | 45.0 | T | 5#3# | 0 | 68.4 | 130 | 65 | 6.02 | 119.42 |
| | | | B _a | 68.4 | | | | | | |
| 200 | 51.0 | T | 5#4# | 0 | 77.0 | 78 | 43 | 5.17 | 133.17 | |
| | | B _a | 75.1 | | | | | | | |
| Doppler Frequency (Hz) | | | 505 | | | Velocity of Target (cm/s) | | | 10 | |
| | | | | | | | | | | |