READ THE MANUAL BEFORE OPERATING THIS PRODUCT

COMPEX

FUSE™

Lidocaine TENS with up to 4% Lidocaine

Version: A0 (2019-6-6)

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INTRODUCTION

Compex® Fuse™ helps deliver electric pulses generated to the user's skin through the electrodes. The portable and compact device has multiple modes of different pulse frequencies, covering Transcutaneous Electrical Nerve Stimulation (TENS) and Powered Muscle Stimulation (PMS) that is also called Electrical Muscle Stimulation (EMS) for over-the-counter use. It includes operating elements of ON/OFF button, intensity increase/mode selection button, and intensity decrease/timer selection button, and could be attached and detached to the electrode through the connectors or the lead wire. When the lead wire is used, one end of the lead wire is connected to the device, and the other end of the lead wire is connected to the electrode pad.

INDICATIONS FOR USE

OVER-THE-COUNTER USE:

TENS:

To be used to aid the temporary relief of pain associated with sore and aching muscles in the shoulder, waist, back, arm, and leg, due to strain from exercise or normal household and work activities.

It is also intended to aid the symptomatic relief and management of chronic, intractable pain and relief of pain associated with arthritis.

PMS:

To stimulate healthy muscles in order to help improve and facilitate muscle performance. To be used to help the improvement of muscle tone and firmness, and for strengthening muscles in the arms, abdomen, legs, and buttocks. Not intended for use in any therapy or for the treatment of any medical conditions or diseases.

It is also intended to help temporarily increase local blood circulation in the healthy muscles of lower extremities.

SAFETY WARNING

CAUTION, CONSULT ACCOMPANYING DOCUMENTS

CONTRAINDICATIONS

- Do not use this device on persons who have a cardiac pacemaker, implanted defibrillator, or other implanted metallic or electronic device, because this may cause electric shock, burns, electrical interference, or death.
- Do not use this device on persons whose pain syndromes are undiagnosed.

WARNINGS

WARNING: Use carefully as directed and for external use only. May cause serious burns. Do not use over sensitive skin areas or in the presence of poor circulation. The unattended use of this device by children or incapacitated persons may be dangerous. To reduce the risk of buns, electric shock, and fire, this device must be used in accordance with the instructions.

- Do not crush the device and its electrodes, and avoid sharp folds.
- Carefully examine the device and its electrodes, and do not use if they show any sign of deterioration.
- Do not tamper with this device and its electrodes in any way. There are no user serviceable parts. If for any reason they do not function satisfactorily, return to the authorized service center at address given.
- The long-term effects of chronic electrical stimulation are unknown.
- Stimulation should not be applied over the carotid sinus nerves, particularly in persons with a known sensitivity to the carotid sinus reflex. Carotid sinus is located on both sides of the neck (Refer to the page of Recommended Use Positions for details).

- Stimulation should not be applied over the front neck or mouth. Severe spasm of the laryngeal and pharyngeal muscles may occur and the contractions may be strong enough to close the airway or cause difficulty in breathing. Stimulation over the neck can also cause adverse effects on heart rhythm or blood pressure.
- Do not apply stimulation across the patient's chest, because the introduction of electrical current into the chest may cause rhythm disturbances to the patient's heart, which could be lethal.
- Do not allow contact with the eyes.
- Since the effects of stimulation of the brain are unknown, stimulation should not be applied across the head, and electrodes should not be placed on opposite sides of head.
- Stimulation should not be applied over swollen, infected, or inflamed areas or skin eruptions, e.g., phlebitis, thrombophlebitis, varicose veins, etc.
- Stimulation should not be applied over, or in proximity to, cancerous lesions.
- Do not use at the same time as other topical analgesics.

PRECAUTIONS

- Safety of stimulation use during pregnancy and breastfeeding has not been established. Consult with your physician before use during pregnancy and breastfeeding.
- Caution should be used for persons with suspected or diagnosed heart problems.
- Caution should be used for persons with suspected or diagnosed epilepsy.
- Caution should be used if you have any of the following:
  - if you have a tendency to bleed internally following an injury;
  - if you recently had surgery, or have ever had surgery on your back;
  - if areas of skin lack normal sensations, such as skin that is numb.
- Consult with your physician before use over the menstrual uterus.

- You may experience skin irritation or hypersensitivity due to the electrical stimulation or electrical conductive medium. The irritation can usually be reduced by using an alternate conductive medium, or alternate electrode placement.
- Do not use this device while driving, operating machinery, or during any activity in which involuntary muscle contractions may put the user at undue risk of injury.
- Keep this device and electrodes out of reach of children and pets. If swallowed, get medical help or contact a Poison Control Center right away.
- Do not use this device in high humidity areas such as a bathroom.
- Stop using this device at once if you feel discomfort, dizziness or nausea, and consult your physician.
- Do not attempt to move the electrodes while the device is operating.
- Do not apply stimulation of this device in the following conditions:
  - across the chest because the introduction of electrical current into the chest may cause rhythm disturbances to the heart, which could be lethal;
  - over open wounds or rashes, or over swollen, red, infected, or inflamed areas or skin eruptions (e.g., phlebitis, thrombophlebitis, varicose veins);
  - in the presence of electronic monitoring equipment (e.g., cardiac monitors, ECG alarms);
  - on children or incapacitated persons.
- Be aware of the following:
  - consult with your physician before using this device;
  - this device is not effective for pain associated with Central Pain Syndromes, such as headaches;
  - this device is not a substitute for pain medications and other pain management therapies;
  - this device is a symptomatic treatment and, as such, suppresses the sensation of pain that would otherwise serve as a protective mechanism;

  - stop using the device and consult with your physician if the device does not provide pain relief;
  - use this device only with the electrodes and accessories recommended for use by the manufacturer. The electrodes may be packaged together with the device or packaged separately as the replacement.
- Store the device away from high-temperature and direct-sunlight. Storage outside of stated storage temperature may result in measurement error or device malfunction.
- Do not share the use of the electrodes with others; electrodes are intended for single person use.
- This device contains batteries. If overheating of the device occurred, stop the operation immediately and contact customer support.
- Dispose of this battery-containing device according to the local, state, or federal laws. Dispose of used electrodes in manner that always keeps product away from children and pets.
- Skin burns may occur, and check the skin under the electrode periodically.

ADVERSE REACTIONS

- You may experience skin irritation and burns beneath the stimulation electrodes applied to the skin; check skin under electrodes frequently.
- You should stop using the product and should consult with your physician if you experience adverse reactions from the device and/or electrodes.
- You should stop using the device and should consult with your physician if condition worsens, redness is present, irritation develops, symptoms persist for more than 7 days or clear up and occur again within a few days, or you experience signs of skin injury, such as pain, swelling, or blistering where the product was applied.

SYMBOL AND TITLE

Information essential for proper use shall be indicated by using the corresponding symbols. The following symbols may be seen on the device and labeling.


CAUTION, Avoid injury. Read and understand owner's manual before operating this product


Type BF applied part


Consult instructions for use


Caution, consult accompanying documents


Use by date


Date of manufacture

Batch code

Catalogue number

Serial number


Manufacturer


Temperature limitation


Humidity limitation

Non-sterile

MR Unsafe - keep away from magnetic resonance imaging (MRI) equipment

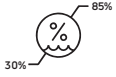

Fragile, handle with care


Keep away from rain


Product packaging is able to be recycled

ENVIRONMENTAL CONDITION FOR TRANSPORT AND STORAGE

Normal working ambient temperature: 10~40°C



Normal working ambient humidity: 30~85%

Store and transport ambient temperature: -10 ~50°C

Store and transport ambient humidity: 30~90%

HOW THE DEVICE WORKS

The device has multiple program modes of TENS and PMS. If you are using the device for the first time, it is recommended that you start with the default Mode 1. Some modes are particularly effective for certain users, but you may need to select both the mode and the electrode position that are the best for you. You may need to try a few positions and modes before finding the one that suits you the best. The best choice of modes and positions may vary from one user to another, in spite of having the same type of symptom.

Program name	Timer (min)	Frequency (Hz)	Pulse Width (μs)
Mode 1 (combination mode): PMS/TENS	30, 40, 50, 60, 10, 20	1.2-160	100
Mode 2: TENS	30, 40, 50, 60, 10, 20	62.5	100
Mode 3: PMS	30, 40, 50, 60, 10, 20	12.5-55.5	100
Mode 4: TENS	30, 40, 50, 60, 10, 20	1.2	100
Mode 5: TENS	30, 40, 50, 60, 10, 20	100	100
Mode 6: TENS	30, 40, 50, 60, 10, 20	160	100

TECHNICAL INFORMATION

Model/type	LidoTENS	Weight	20g
Power supply	Powered by a built-in 3.7V li-ion battery	Automatic shutoff	30 min
Waveform and wave shape	Biphasic rectangular wave pulse	Degree of protection against electric shock	Type BF applied part
Pulse width	100μsec	Type of protection against electric shock	Internally powered equipment
Pulse frequency	1-160Hz (Hz=stimulation times per second)	Grade of waterproof	IP22
Output Voltage	Max. 70Vpp ±20%(at 500Ω)	Product warranty	1 year
Treatment time	30, 40, 50, 60, 10, 20 min	Lifetime for electrode	Storage for 2 years (no use), Times of reusable: 30 times
Output intensity	0 to 20 levels, adjustable	Mode of operation	Continuous operation
Modes	6 auto modes	Software version	A0
Typical operation time of Battery	If to use at the highest level, the battery can be used for about 150 min after fully charged.	The time required for me equipment to warm from the minimum storage temperature between uses until it is ready for intended use	30 min
Behaviour of me equipment while the rechargeable internal electrical power source is charging:	The indication light will flash during charging and will be still with full capacity.	The time required for me equipment to cool from the maximum storage temperature between uses until it is ready for intended use	15 min
Typical service life of Battery	300 times of recharging	Adapter for charging	Please use output DC5V and output current 0.3-2.0A adapter for charging
Note: Not intended to be sterilized.			
Not for use in an OXYGEN RICH ENVIRONMENT			

SETUP

Unpack the box of the product, and take the product and accessories out.

Accessories included in the package

1 TENS unit

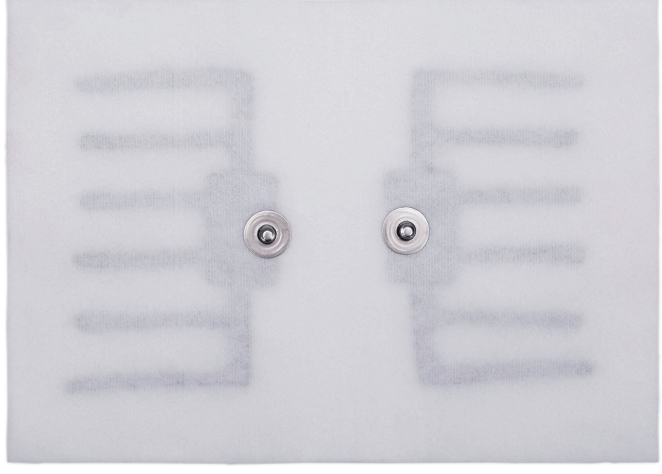
1 electrode patch

1 USB cable

1 Manual

TENS Unit (Side View)

USB Cable



Electrode Pack

OPERATING INSTRUCTION

The following Operating Instructions are used to guide the operation, and the details about each step are listed in the following table.

1st Step – Check the battery power for the device The device comes with a built-in rechargeable battery, and could be used as received. The USB cable included could be used to charge the device. The light indicator on the device is flashing during charging, and becomes solid when the device is charged fully. Note: Use the UL- or ETL- certified adapter of 5V and 0.3-2A for the battery charging.	 Charge Device
2nd Step - Install the electrode onto the device Snap the enclosed electrode onto the device through the contact connectors. This should be done prior to applying the device onto the skin of the treatment area.	 Electrode installation
3rd Step - Put the electrode-installed device on the stimulation-needed body area Carefully remove the protective cover film from the electrode pad/patch, and avoid fingers contacting to the sticky electrode gel. Place the electrode-installed device onto the treatment area. Press down firmly and ensure a full and firm contact to the body area of treatment. **Note:** Keep the skin clean before placing the electrode patch-installed device and wash hands if fingers contact to the sticky electrode gel.	 Place the device on the body area
4th Step - Press the "ON/OFF" to turn on the power Hold the On/Off button to turn on the unit, illustrated by the lit-up indicator on the front.	 On/Off

5th Step - Select one of the stimulation modes Change the output stimulation modes by pressing and holding the + button, and the mode change could be seen on the flashing times of the light indicator. **Note:** When the light indicator flashes once, it means Mode 1; when the light indicator flashes twice, it indicates Mode 2.	 Mode change
6th Step - Choose the stimulation time The default timer is 30 min. Change the timer by pressing and holding the - button, and the timer change could be seen on the flashing times of the light indicator. **Note:** When the light indicator flashes once, it means 10min; when the light indicator flashes twice, it indicates 20min.	 Timer change
7th Step - Adjust the stimulation intensity Press and release the "+" button to increase the stimulation intensity, and press and release the "-" button to decrease the intensity. **Note:** With the increase of intensity, you may experience sensations like tingling, vibration, etc. Therefore, gradually increase the intensity, and stop increasing when a comfortable level is reached.	 Intensity change
8th Step - Stay with the stimulation After the above mode, timer, and intensity are set, the stimulation treatment provided by the device will last until the device turns off.	 Stay with the stimulation treatment
9th Step - Press the "ON/OFF" to turn off the power after done When the timer is up, the device will turn off automatically. The device could be also turned off by holding the On/Off button. The electrode patch/pad could remain on the treatment area for up to 12 hours, while the device turns off automatically or manually. Note: When not in use, store the device and accessory in a cool place, out of direct sunlight.	 On/Off

If using the device for the first time, you may start from the default Mode 1. And if you expect to use the specific mode, please refer to the following for details.

USE AS TENS

The device could be placed directly on the treatment site where you are experiencing pain (such as pain associated with sore and aching muscles and pain associated with arthritis). For example, if the treatment site is on the calf, place the TENS device above the pain site.

Arthritis involves inflammation on joints of the body, usually producing pain and stiffness. The electrode is placed on or near the area of the arthritis pain. The TENS mode of the device generates electrical pulses that are sent via the electrode for pain relief.

USE AS EMS

The device could be placed directly on the body area where you are expecting the muscle strengthening. For example, if the body area is the leg, place the device on the leg site.

RECOMMENDED PRACTICE FOR BOTH TENS AND EMS:

- Start from the lowest intensity and gradually adjust the intensity to a comfortable level. Duration is the preset timer (such as 30 minutes) for each skin area. Frequency is 2 times per day per skin area. The electrode patch/pad could remain on the treatment area for up to 12 hours per day, while the device is turned off automatically or manually. Consult with your physician for longer and more frequent uses.
- Good skin care is important for a comfortable use of device. Be sure the treatment site is clean of dirt and body lotion.
- Keeping the electrode in the storage bag after use will extend its lifespan. The electrode pad is disposable and should be replaced when it loses the adhesiveness. To purchase additional electrodes, please contact the seller.

The exact percentage of ingredients used in the electrode patch/pad is withheld as the trade secret and may be disclosed as requested. Electrodes with a shelf life of 2 years are intended for single person use, and could be reused and replaced. If the electrode no longer contacts well with your body skin, it is time to replace it. If needed, you could use a damp and clean cloth to wipe electrodes between uses. Please see the following for the Use Direction, Removal and Storage of the electrode used for both TENS and EMS.

USE DIRECTION

1. Clean skin thoroughly prior to each application of electrodes, which will not contact well if any lotion, make-up, or dirt is left on the body skin.
2. Ensure the device is off before applying the electrode to it.
3. Apply the electrode-installed device firmly to the body skin.

REMOVAL AND STORAGE

1. Turn the device off before removing the electrode from the body skin.
2. Grab the edge of the electrode and remove from the body area.
3. When not in use and/or between each use, store the electrode in the re-sealable bag, out of direct sunlight and out of reach of children and pets.

PERFORMANCE SPECIFICATIONS

Power Source	3.7V Battery
Number of Output Modes	6 preset modes
Timer Range (minutes)	10-60
Dimensions (mm) [L x W x D]	52 x 44 x 12 mm
Waveform	Biphasic
Shape	Rectangular
Maximum Output Voltage	70V@500Ω
Maximum Output Current	80mA@500Ω
Maximum Pulse Duration	100μSec
Maximum Frequency	160Hz

CLEANING AND MAINTENANCE

Please use wipe to clean the device first, and then use the dry cloth to wipe it again. The electrodes coming with the device are disposable, and should be replaced when their adhesiveness becomes worse. Contact the seller for replacements.

TROUBLE SHOOTING

If your device is not operating properly, please check below for common problems and suggested solutions. If the recommended action does not solve the problem, please contact the seller.

STIMULATION IS WEAK OR NON-EXISTENT

- Be sure skin is clean and pads are firmly attached to skin.
- The battery is low and needs to be charged.

DEVICE DOES NOT TURN ON

- Check if battery is low and needs to be charged.

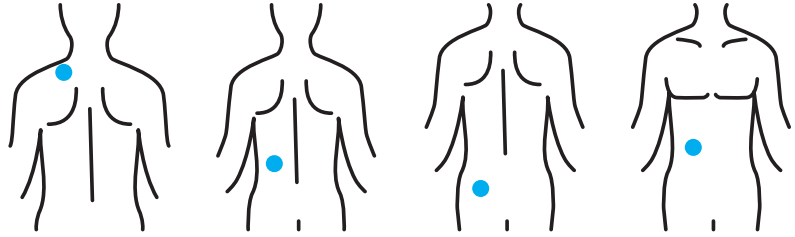
ELECTRODES ARE NOT STICKY

- The pads will lose adhesiveness over use. Please contact the distributor or dealer for replacements.

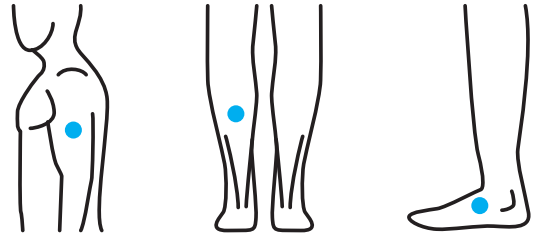
SKIN TURNS RED

- Do not use the product if skin irritation or redness occurs.
- If problem persists, contact your physician.

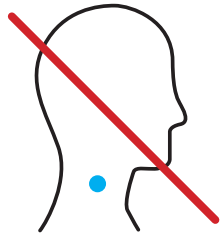
RECOMMENDED USE POSITIONS



SHOULDERS **BACK** **BUTTOCKS** **ABDOMEN**



ARMS **LEGS** **FEET**



NEVER APPLY ELECTRODES ON THE THROAT OR BOTH SIDES OF THE NECK, WHERE THE CAROTID SINUS NERVES ARE LOCATED.

WARRANTY

This device carries a limited warranty of one year from the date of delivery. The warranty applies to the device only, and the accessories are not covered by this warranty.

During the warranty period, defective items will be repaired or replaced at no charge. Any evidence of misuse, abuse, alternations, or externally caused damage may have this warranty invalid.

For more information, please contact the manufacturer.

ELECTROMAGNETIC COMPATIBILITY AND FCC COMPLIANCE STATEMENT

1. This product needs special precautions regarding electromagnetic compatibility (EMC) and needs to be installed and put into service according to the EMC information provided, and this unit can be affected by portable and mobile radio frequency (RF) communications equipment.
2. Do not use a mobile phone or other devices that emit electromagnetic fields, near the unit. This may result in incorrect operation of the unit.
3. **CAUTION:** This unit has been thoroughly tested and inspected to assure proper performance and operation!
4. **CAUTION:** This machine should not be used adjacent to or stacked with other equipment and that if adjacent or stacked use is necessary, this machine should be observed to verify normal operation in the configuration in which it will be used.

NO ESSENTIAL PERFORMANCE WAS IDENTIFIED.

GUIDANCE AND MANUFACTURE'S DECLARATION – ELECTROMAGNETIC EMISSION		
The device *is* intended for use in the electromagnetic environment specified below. The customer of the user of the device should assure that it is used in such an environment.		
EMISSION TEST	**COMPLIANCE**	**ELECTROMAGNETIC ENVIRONMENT – GUIDANCE**
RF emissions CISPR 11	Group 1	The device 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 CISPR 11	Class B	The device 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 61000-3-2	Class A	
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Complies	

GUIDANCE AND MANUFACTURE'S DECLARATION – ELECTROMAGNETIC IMMUNITY			
The device 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	±8 kV contact ±15 kV air	±8 kV contact ±15 kV 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 61000-4-4	±2 kV for power supply lines ±1 kV for input/output lines	±2kV 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 differential mode	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% UT (>95% dip in UT) for 0.5 cycle 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 5 sec	<5% UT (>95% dip in UT) for 0.5 cycle 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 5 sec	Mains power quality should be that of a typical commercial or hospital environment. If the user of the device requires continued operation during power mains interruptions, it is recommended that the device be powered from an uninterruptible power supply or a battery.
Power frequency (50Hz/60Hz) 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.
NOTE UT is the a.c. mains voltage prior to application of the test level.			

GUIDANCE AND MANUFACTURE'S DECLARATION – ELECTROMAGNETIC IMMUNITY			
The device 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 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 device, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance $d = 1{,}2\sqrt{P}$
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz	3 V/m	$d = 1{,}2\sqrt{P}$ 80 MHz to 800 MHz $d = 2{,}3\sqrt{P}$ 800 MHz to 2,5 GHz 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). 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: 
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 device is used exceeds the applicable RF compliance level above, the device should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the device.

b Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.

RECOMMENDED SEPARATION DISTANCES BETWEEN PORTABLE AND MOBILE RF COMMUNICATIONS EQUIPMENT AND THE DEVICE.

The device is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the device can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the device 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	80 MHZ TO 800 MHZ	800 MHZ TO 2.5 GHZ
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
1.0	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 metres (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.

FCC COMPLIANCE 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.

The subject device 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.

The product generates, uses, and can radiate radio frequency energy and, if not installed and used accordance with the instructions, may cause harmful interference to radio communications.

However, there is no guarantee that the interference will not occur in a particular installation. If the product does cause harmful interference to radio or television reception, which can be determined by turning the product on or off, the user is encouraged to try to correct the interference by one or more of the following measures:

a. Reorient or relocate the receiving antenna;

b. Increase the separation between the product and the receiver;

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

d. Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.

Changes or modifications to this product not expressly approved by the party responsible for compliance could void the user's authority to operate the equipment.

CONTACT INFORMATION

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