	USER MANUAL	WARNINGS PRECAUTIONS		CARE AND STORAGE OF ELECTRODE PADS	TROUBLESHOOTING GUIDE		ELECTROMAGNETIC COMPATIBILITY (EMC)
	INTRODUCTION	 If you are in the care of a physician, consult your physician before using this device. 	The long-term effects of TENS therapy (cutaneous electrodes for electrical stimulation) are unknown.	Clean the gel surface of the electrode pad by holding under dripping water and gently rubbing with your fingers for 5-10 seconds. Allow	Problem	Possible Cause Corrective Acti	Medical Electrical Equipment needs special precautions regarding EMC and needs to be installed and put into service according to the
	What is TENS? TENS stands for Transcutaneous Electrical Nerve Stimulation. TENS therapy uses mild electrical impulses applied to the skin for pain relief. The SmartRelief device is a battery-powered TENS device that transmits a	 If you have had medical treatment for your pain, consult with your physician before using this device. If your pain does not improve or becomes more than acute, stop using the device and consult with your physician. Do not use this device if you have a cardiac pacemaker, implanted 	TENS is not a substitute for pain medications and other pain management therapies. Use caution if electrode pads are applied over areas of skin that lack normal sensation. Replace self-adhesive electrode pads if they no longer stick firmly to	to air dry. This should only be done when the electrode pad has lost adhesion. • Place electrode pad on the clear protective sheet and store the electrode pad in its re-sealable bag after each use. The life of the electrode pads varies depending on skin conditions, skin preparation,	No stimulation felt	Dead battery Battery installed incorrectly Dry electrode Electrode torn or broken Replace battery (CR 2032) Insert battery correctly Replace electron	EMC information provided in this manual. Portable and mobile Radio Frequency (RF) communications equipment can affect Medical Electrical Equipment.
ICYHOT®	mild electrical impulse, through conductive electrode pads applied on the skin, to underlying nerve fibers and muscles. TENS therapy is a safe and effective drug-free method of reducing pain that is used by physicians and	defibrillator, or other implanted metallic or electronic device. Doing so could cause electric shock, burns, electrical interference, or death. Do not place this device on your head.	vour skin. Reuse of the electrode pad by another user could cause the cross-infection of skin diseases. Clean and dry the area to which you will apply the electrode pad with	storage and climate.	Intermittent stimulation	Poor connection Damaged electrode Re-attach to electrode Replace electrone	The use of accessories, transducers and cables other than those specified by the manufacturer may result in increased emissions or decreased immunity of the SmartRelief device.
SMARTRELIEF™ TENS THERAPY	pain clinics around the world. INDICATIONS FOR USE To be used for temporary relief of pain associated with sore and aching	wat was a borner of place this device over your rieck. Doing so could cause severe muscle spasms resulting in closure of your airway, difficulty in breathing, or adverse effects on heart rhythm or blood pressure. Do not place this device across your chest. The introduction of an electrical current into the chest may cause rhythm disturbances to your	water or alcohol prior to application. Use of accessories not approved by the manufacturer may cause harm or injury. Operation in close proximity to short wave or microwave therapy equipment may produce instability in the stimulator output.	On/Off Switch On Indicator (LED) Increase Button Decrease Button	Stimulation not powerful enough	Electrode dry or damaged Low battery Poor electrode contact with skin Replace electrode Replace battery Clean skin and replace electrod	with other equipment. If adjacent or stacked use is necessary, the SmartRelief device should be observed to verify normal operation
	muscles due to strain from exercise or normal household and work activities. To be used for the symptomatic relief and management of chronic, intractable pain and relief of pain associated with arthritis.	 beart, which could be lethal. Do not use this device in the presence of cardiac pacemakers or implants. Do not use this device during pregnancy. Do not place this device over the carotid sinus nerves, the front of the neck, or around the mouth. 	Keep the SmartRelief power unit clean by wiping with a damp cloth. Do not immerse. Do not disassemble the SmartRelief power unit. Electrode pads should only be applied to normal, intact, clean, healthy skin. The size, shape, and type of electrodes may affect the effectiveness of pain relief. Use only SmartRelief electrodes with your SmartRelief device		Stimulation surges	Electrode not staying attached to skin Re-attach elect to power unit Clean skin and re-attach electre Replace electro	l l
Relief for Muscle Aches and Pains, Arthritis, and Chronic Pain	CONTRAINDICATIONS Do not use this device if you have a cardiac pacemaker, implanted defibrillator, or other implanted metallic or electronic device. Such use could cause electric shock, burns, electrical interference, or death.	 Do not place this device over open wounds, sores or rashes, or over swollen, red, infected, or inflamed areas or skin eruptions (e.g., phlebitis, thrombophlebitis, varicose veins). Do not use this device over, or in proximity to, cancerous lesions. Do not use this device when in the bath or shower. 	and only use as directed. The adhesive characteristics of electrode pads may affect the safety and effectiveness of electrical stimulation. Clean and/or replace your electrode pads as directed. Using electrode pads that are too small or no longer adhere could result	3V Battery CR 2032 Battery Drawer	Unwanted muscle twitches Stimulation	Intensity too high Electrode placement Hove electrode Bectrode placement Hove electrode Hove electrode	
Visit www.smartrelief.com for more information		 Do not use this device while sleeping. Do not use this device while driving, operating machinery, or during any activity in which electrical stimulation can put you at risk of injury. Do not use this device in the presence of electromagnetic fields. 	in discomfort, irritation, or minor skin burns. Choking hazard: Do not swallow battery. Keep this device out of reach of children.	i i	ineffective against pain Stimulation feels weaker a few	Intensity too low Increase intens Normal operation – electrical pattern is cyclic desired	_ 1
on SmartRelief and coupon offers.	instructions.	 Do not use this device if case has been damaged. Do not use this device if you don't know the source of your pain. Do not use this device on children. 	ADVERSE REACTIONS		LED light not lit	Dead battery Replace battery	1
 	Distributed by Chattem, Inc. P.O. Box 2219 Chattanooga, TN 37409-0219 USA		If you experience adverse reactions, stop using this device and consult with your physician. Users with sensitive skin may experience skin irritation in the area where the electrode pad is applied. You may experience headache and other painful sensations during or following the application of electrical stimulation near your eyes and to	BACK I Patent Pending	by temporary irrita	s experience skin sensitivity to TENS characterization or itching. Use at a lower intensity until you amfortable with the TENS sensation.	
0075688-01	2	3	your head and face. 4	5			6 7

Electromagnetic Immunity The SmartRelief device is intended for use in the electromagnetic environment specified below. The customer or the user of the SmartRelief device should assure that it is used in such an environment

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hospital environment.

hospital environment.

Electromagnetic Environment - Guidance

Mains power quality should be that of a typical commercial or

Mains power quality should be that of a typical commercial or

Mains power quality should be that of a typical commercial or

recommended that the SmartRelief device be powered from an

Power frequency magnetic fields should be at levels characteristic

of a typical location in a typical commercial or hospital environment.

continued operation during power mains interruptions, it is

uninterruptible power supply or a battery.

hospital environment. If the user of the SmartRelief device requires

Immunity Test IEC 60601 Test Level Compliance Level Electrostatic +/- 6 kV contact +/- 6 kV contact discharge +/- 8 kV air +/- 8 kV air IEC 61000-4-2 Electrical fast +/- 2 kV for power Not Applicable supply lines transient / burst IEC 61000-4-4 +/- 1 kV for input/ output lines Surge Not Applicable +/- 1 kV line(s) to IEC 61000-4-5 +/- 2 kV line(s) to earth <5% Ur Not Applicable Voltage dips, short (>95% dip in Ur) interruptions and for 0.5 cycle voltage variations or power supply input 40% Ur (60% dip in Ur) for 5 cycles IEC 61000-4-11 70% Ur (30% dip in Ur) for 25 cycles <5% Ur (>95% dip in Ur) for 5 seconds 3 A / m Power frequency 3 A / m magnetic field

IEC 61000-4-8

NOTE Ur is the A.C. mains voltage prior to application of the test level.

Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.

from structures, objects and people.

should be less than 3 V/m.

Immunity Test

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 SmartRelief device is used exceeds the applicable RF compliance level above. The SmartRelief device should be observed to verify normal operation. If abnormal performance is observed additional measures may be necessary such as reorienting or relocating the SmartRelief device.

b) Over the frequency range 150 kHz to 80 MHz, field strengths

The SmartRelief device is intended for use in the electromagnetic environment specified below. The customer or the user of the SmartRelief device should assure that it is used in such an environment. IEC 60601 Test Level Compliance Level Electromagnetic Environment - Guidance

		Compilation Lotte
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80MHz	3 Vrms
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.5GHz	3 V/m
	10 V/m 26MHz to 1000MHz	10 V/m

Guidance and Manufacturer's Declaration

Electromagnetic Immunity

NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection

> 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:

 $d = 1.2\sqrt{P}$

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Portable and mobile RF communications equipment should be used no

closer to any part of the SmartRelief device, including cables, than the

recommended separation distance calculated from the equation

Where P is the maximum output power rating of the transmitter in

watts (W) according to the transmitter manufacturer and d is the

Field strengths from fixed RF transmitters as determined by an

recommended separation distance in meters (m).

applicable to the frequency of the transmitter.

Recommended separation distance

 $d = 1.2\sqrt{P}$ 80 MHz to 800 MHz

 $d = 2.3\sqrt{P}$ 800 MHz to 2.5 GHz

The SmartRelief device is intended for use in the electromagnetic

flicker emissions

IEC 61000-3-3

Electromagnetic Emissions

environment specified below. The customer or the user of the SmartRelief device should assure that it is used in such an environment. Emissions Test Compliance Electromagnetic Environment - Guidance RF emissions Group 1 The SmartRelief device uses RF energy only for its interna CISPR 11 function. Therefore its RF

Guidance and Manufacturer's Declaration -

emissions are very low and are not likely to cause any interference in nearby electronic equipment RF emissions The SmartRelief device is suitable for use in all CISPR 11 establishments, including domestic establishments and Harmonic Not Applicable those directly connected to the emissions public low-voltage power IEC 61000-3-2 supply network that supplies buildings used for domestic Voltage Not Applicable fluctuations /

NOTE 1 At 80 MHz and 800 MHz, the separation distance for the higher

Electromagnetic propagation is affected by absorption and

frequency range applies.

NOTE 2 These guidelines may not apply in all situations.

reflection from structures, objects and people

portable and mobile RF communication equipment Store between: and the SmartRelief device The SmartRelief device is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user

of the SmartRelief device can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the SmartRelief device as recommended below, according to the maximum output power of the communications equipment.

Recommended separation distances between

nal I	Rated maximum	Separation distance according to frequency of transmitter m			Intensity Control: Adjustable in 63 sto Timer: Device turns off after 30 minu	
	output power of transmitter	150 kHz to 80 MHz d = 1.2 √ P	80 MHz to 800 MHz d = 1.2 √ P	800 MHz to 2.5 GHz d = 2.3 √ P	Waveform: Asymmetrical Biphasic Pu For Use in Home, Office or Workplace Operating Temperature: 10 - 40 degree Operating Humidity: 30% to 75% Atmospheric Pressure 50 - 106 kPa	
	0,01	0.12	0.12	0.23	I	
ıd	0,1	0.38	0.38	0.73	1-100 PPS	
the I	1	1.2	1.2	2.3	1	
	10	3.8	3.8	7.3	! <u>Г</u> і	
	100	12	12	23		
	For transmitters rat	ed at a maximum	output power not	listed above, the	0.014 - 0.031 Vr	

recommended separation distance d in meters (m) can be estimated

P is the maximum output power rating of the transmitter in watts (W)

according to the transmitter manufacturer.

using the equation applicable to the frequency of the transmitter, where

-40 and 70 degrees C

10% - 90% Relative humidity

Atmospheric Pressure 50 - 106 kPa

ENVIRONMENTAL AND TECHNICAL

Dimensions: 64x38x13mm Weight: 20g

Power Supply: 3V Battery CR 2032 x 1

Channels: 1 channel output through female snaps Output Current: 0-63mA peak into a 500-ohm load Intensity Control: Adjustable in 63 steps

Timer: Device turns off after 30 minutes Waveform: Asymmetrical Biphasic Pulse, 0 net DC Charge For Use in Home. Office or Workplace Operating Temperature: 10 - 40 degrees C



