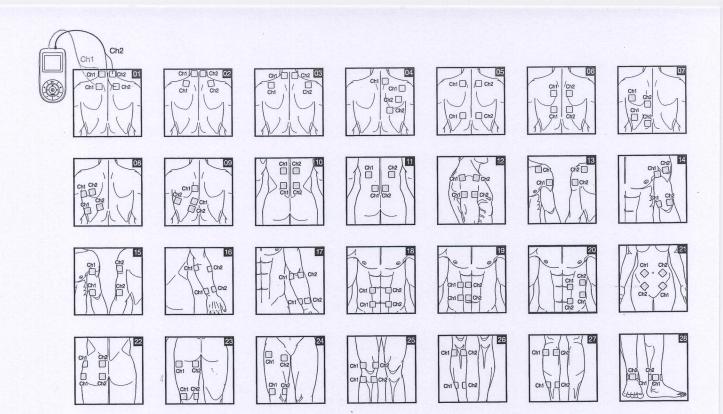


TENS Digital Pain Relief unit Instruction for Use





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Please read these instructions for use carefully and keep them for later use, be sure to make them accessible to other users and observe the information they contain

1. Getting to know your machine

1.1 What is Digital EMS/TENS and what can it do?

Digital EMS/TENS belongs to the group of electrical stimulation systems. It has three basic functions, which can be used in combination:

- 1. Electrical stimulation of nerve tracts (TENS)
- Electrical stimulation of muscle tissue (EMS).
- 3. A massage effect generated by electrical signals.

For this, the device has two independent stimulation channels and four self-adhesive electrodes. It has various versatile functions in order to increase general wellbeing, alleviate pain, maintain physical fitness, relaxation, revitalise muscles and combat fatigue. You can either select these functions from preset programs or determine them yourself according to your requirements. The operating principle of electrical stimulation equipment is based on simulating the body's own pulses which are transmitted transcutaneously to nerve or muscle fibres by means of electrodes. The electrodes can be attached to many parts of the body, whereby the electrical stimuli are safe and practically painless. You merely feel a gentle prickling or vibrating. The electrical pulses transmitted to the tissue affect the transmission of stimulation in nerve conductions as well as neural nodes and muscle groups in the field of application.

The effect of electrical stimulation is generally only recognisable after regularly repeated application. Electrical stimulation does not replace regular exercising of the muscle, but is able to reasonably supplement the effect.

TENS, transcutaneous electrical nerve stimulation, means electrical stimulation of nerves through the skin. TENS is recognised as a clinically proven, effective, non-medication method of treating pain from certain causes. It is free from side-effects when used properly, and can also be used as a simple means of self-treatment. The pain-alleviating or suppressing effect is also achieved by preventing pain from being passed on into nerve fibres (above all through high-frequency pulses) and increasing the secretion of the body's own endorphins which reduce awareness of pain through their effect on the central nervous system. The method is scientifically underpinned and medically approved.

Any symptoms indicating TENS application must be clarified with the doctor

in charge of your treatment. The latter will also give you information on the respective benefits of TENS self-treatment.

TENS is clinically tested and approved with the following applications:

- Back pain, particularly also lumbar and cervical spine problems · Joint pain (e.g. knee joint, hip joint, shoulder)
- Neuralgia
- Women's period pains
- · Pain after injuries to musculoskeletal system
- Pain with circulatory problems
- · Chronic pain through various causes

Electrical muscle stimulation (EMS) is a widespread and generally recognised method and has been used for years in sports and rehabilitation medicine In the sports and fitness field, one of the uses of EMS is as a supplement to conventional muscle training, in order to increase the efficiency of muscle groups and adapt physical proportions to the desired aesthetic results. EMS application goes in two directions. On the one hand, a targeted strengthening of musculature can be produced (activating application) and on the other hand a relaxing, resting effect can also be achieved (relaxing application).

Active application includes the following:

- Muscle training to improve endurance performance and/or
 Muscle training to support the strengthening of certain muscles or muscle groups in order to achieve desired changes to body proportions.

Relaxing application includes the following:

- Muscle relaxation in order to loosen up muscular tension
- Improving muscular fatigue symptoms
- · Accelerating muscle regeneration after high muscular performance (e.g. after a marathon)

Through **integrated massage technology**, Digital EMS/TENS also offers the possibility of reducing muscular tension and combating fatigue symptoms using a program based on real massage in terms of sensation and effect.

Using the positioning suggestions and program charts in this manual, you can determine the machine setting fast and easily for the respective application (depending on the region of the body affected) in order to ensure the desired

Through the two separately adjustable channels, the Digital EMS/TENS has the advantage of adapting the intensity of the pulses independently of each other

to two parts of the body to be treated, for example in order to cover both sides of the body or ensure even stimulation of larger areas of tissue. The individual intensity setting of each channel also allows you to treat two different parts of the body at the same time, whereby it is possible to save time compared with individual sequential treatment.

1.2 Contents

- 1x Digital EMS/TENS machine (incl. belt clip)
- 2x connecting cable 4x adhesive electrodes (45 x 45 mm)
- 3x AAA batteries
- Storage pouch
- these instructions for use

Replacement Pads

8x adhesive electrodes (45 x 45 mm), item code 1297708

(i) 2. Important Information

Use of the machine does not replace medical consultation and treatment. In the event of any type of pain or illness, you must therefore always first ask your doctor.

/ WARNING!

In order to prevent any damage to health, we would urgently advise against using the Digital EMS/TENS in the following cases:

- With any implanted electrical devices (such as pacemakers)
- If there are any metal implants
- When using an insulin pump
- In case of high fever (e.g. >39 °C)
- In case of known or acute cardiac arrhythmias and other disorders in stimulus formation and conduction of the heart
- If suffering from fits (e.g. epilepsy)
- If pregnant
- In case of cancer
- After operations where increased muscle contractions might impair the healing process



- Never use near the heart. Stimulation electrodes should never be placed anywhere on the front of the thorax (marked by ribs and breastbone), but above all not on the two large pectoral muscles. Here it can increase the risk of ventricular fibrillation and lead to cardiac arrest
- On the bony part of the cranium, near the mouth, throat or larynx
- Near the throat/carotid artery
- Near the genitals
- On acutely or chronically diseased (injured or inflamed) skin (e.g. in the event of painful and painless inflammation, reddening, skin rashes (e.g. allergies), burns, bruises, swellings, open wounds and wounds in the process of healing, on operation scars in the process of healing)
- In environments with high humidity such as in the bathroom or when having a
- Do not use after consuming alcohol
- When a high frequency surgical device is connected at the same time

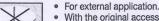
Before using the machine, you should consult with the doctor in charge of your treatment in the event of the following:

- Acute diseases, especially if hypertension is suspected or actually exists, or blood-clotting disorders, tendency to thromboembolic diseases or in case of malignant new masses
- All skin diseases
- Chronic pain disorders which have not been clarified, regardless of the region of the body
- Diabetes
- All sensitivity disorders with reduced pain awareness (such as metabolic disturbances)
- Medical treatments carried out simultaneously
- Complaints arising as a result of the stimulation treatment
- Constant skin irritation as a result of prolonged stimulation at the same electrode point

riangle important!

Use the Digital EMS/TENS solely:

- On humans
- For the purpose for which it was developed and in the manner specified in these instructions for use. Any improper use can be hazardous.



With the original accessory parts which are supplied and can be re-ordered, otherwise the warranty becomes null and void.

PRECAUTIONARY MEASURES:

Always remove the electrodes from the skin with a moderate pull in order to avoid injury in the event of highly sensitive skin.

Keep the machine away from heat sources and never use it near (~1 m) short-wave or microwave equipment (e.g. mobile phones), as this may lead to unpleasant current spikes.

Do not expose the machine to direct sunlight or high temperatures.

 Protect the machine from dust, dirt and humidity. Never submerge the device in water or other liquids.

The machine is suitable for personal use.

For reasons of hygiene, the electrodes should only be used on one person.

If the machine is not functioning properly, or if you start feeling unwell or there is any pain, stop the application immediately.

To remove or move the electrodes, first switch off the machine or the

appropriate channel in order to avoid unwanted irritation. Never modify any electrodes (e.g. by cutting). This leads to a higher current density and can be dangerous (max. recommended output value for electrodes: 9 mA/cm², an effective current density over 2 mA/cm² requires greater attention).

· Do not use when asleep, when driving a vehicle or at the same time as operating machinery.

Never use with any activities where an unforeseen reaction (e.g. increased muscle contraction despite low intensity) may be dangerous.

Make sure that no metallic objects such as belt buckles or necklaces can come in contact with the electrodes during stimulation. If you wear any jewellery or piercings (e.g. belly piercing) near the application site, you must remove these before using the machine as burning may otherwise occur at some points.

 Keep the device away from children to prevent any hazards.
 Do not confuse the electrode cables and contacts with your headphones or other devices, and do not connect the electrodes to other devices.

· Do not use this machine at the same time as other equipment which sends electrical pulses to your body.

Do not use it near any highly flammable substances, gases or explosives.

Never use accumulators, always use the same types of battery.



 In the initial minutes, carry out the application sitting or lying down in order to avoid any unnecessary risk of injury in the rare event of a vagal reaction (feeling of weakness). If you start feeling weak, switch the machine off immediately and put your legs up (approx. 5–10 minutes).

 It is not advisable to apply any enriching creams or ointments to the skin beforehand, as this greatly increases electrode wear and unpleasant current spikes may occur here.

Damage

- If the device is damaged, do not use it and contact your local Boots store.
- Check the device at regular intervals for signs of wear or damage. If you find any such signs or if the device was used improperly, take it to your local Boots store.

Switch off the device immediately if it is defective or if is not working properly.

- Do not, under any circumstances, attempt to open or repair the machine yourself! Only have repairs carried out by the customer service department or an authorized dealer. If these instructions are not heeded, the warranty becomes null and void.
- The manufacturer assumes no liability for damage caused through improper or wrong use.

3. Current parameters

Electrical stimulation machines operate with the following current settings, which will alter the stimulation effect depending on the setting.

3.1 Pulse shape

This describes the time function of the excitation current whereby monophase pulse trains are differentiated from biphase. With monophase pulse trains, the current flows in one direction. With biphase pulses, the excitation current alternates its direction.

In the Digital EMS/TENS unit, there are only biphase pulse

In the Digital EMS/TENS unit, there are only biphase pulse trains, as they reduce the strain on the muscle, leading to less muscle fatigue as well as safer application.

3.2 Pulse frequency

Frequency indicates the number of individual pulses per second, and is indicated in Hz (Hertz). It can be calculated by working out the inverse value of the periodic time. The respective frequency determines which types of muscle fibre





preferably react. Slow-response fibres tend to react to lower pulse frequencies up to 15 Hz, while fast-response fibres only respond to frequencies over approx. 35 Hz.

With pulses of approx. 45–70 Hz, there is permanent tension in the muscle combined with premature muscle fatigue. Higher pulse frequencies can therefore preferably be used for elasticity and maximum strength training.

3.3 Pulse width

Pulse width is used to indicate the duration of an individual pulse in microseconds. Pulse width also determines the penetration depth of the current whereby in general, the following applies: a greater muscle mass requires a greater pulse width.



3.4 Pulse intensity

Setting the degree of intensity is individually dependent on the subjective feeling of each individual user and is determined by a number of parameters such as application site, skin circulation, skin thickness as well as quality of



site, said circulation, said minimized as well as quality of the electrode contact. The actual setting should be effective but should never produce any unpleasant sensations such as pain at the site of application. While a slight tingling sensation indicates sufficient stimulation energy, any setting which leads to pain must be avoided.

With prolonged application, readjustment may be necessary due to time adjustment processes at the site of application.

3.5 Cycle-controlled pulse parameter variation

In many cases it is necessary to cover the entirety of tissue structures at the site of application by using several pulse parameters. With the Digital EMS/TENS unit, this is done by the existing programs automatically making a cyclical pulse parameter change. This also prevents individual muscle groups at the site of application from getting tired.

With the Digital EMS/TENS unit there are useful presettings for current parameters. With this, you can change the impulse intensity at any time during use. For 6 programs you can also set various parameters for stimulation yourself.

4. Controls

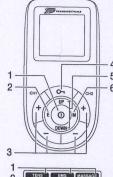
Buttons:

ON/OFF button ①

2 E button (Enter)
3 Intensity setting buttons (Cn1 +/- left, Cn2 +/- right)

4 Keylock On 5 Selection button UP and DOWN

6 M button (Menu)



Display (full screen):

1 Menu TENS / EMS / MASSAGE 2 Low battery

 Keylock
 Timer function (remaining time display) or working time

5 Frequency display (Hz), pulse width (µs) or pause time

Impulse intensity channel 1 (Ch1)
Impulse intensity channel 2 (Ch2)

8 Operating status display
9 Program number

Accessories:

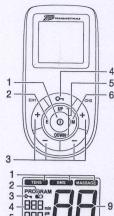
2x connection cable

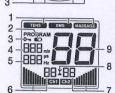
 4x adhesive electrodes (45 x 45 mm)

1x belt clip



Every time a button is pressed, this is acknowledged by a signal tone in order to ensure that any unintentional pressing of a button is detected. This signal tone cannot be switched off.





Keylock O—
Locks the buttons to avoid them being pressed unintentionally.

(1) To activate the keylock, hold down the O— button until the O— symbol is visible in the display (approx. 3 seconds). (2) To deactivate the keylock, hold down the O button until the O symbol disappears from the display (approx. 3 seconds). 5. Start-up

ON/OFF ①

UP and DOWN

M (Menu)

time and pause time.

(1) Menu selection.

Setting pulse intensity.

1. Remove the belt clip from the machine, if attached.

Intensity setting buttons (Chi +/- left, Ch2 +/- right)

2. Press the battery compartment cover on the rear of the device

(1) Press briefly to switch the device on.
(2) Press once to interrupt stimulation treatment.
(3) Hold ON/OFF button ① down (approx. 2 seconds) to switch off the

(1) Navigation between TENS, EMS and MASSAGE menus.
(2) Return to program selection screen or menu selection.

Choice of treatment program, treatment time, frequency, pulse width, working

(2) To confirm a selection made with UP/DOWN, apart from channel intensity.

and slide it downwards.
Insert three AAA 1.5V alkaline batteries. Make absolutely sure that you insert the batteries with the correct polarity as

4. Replace the battery cover carefully (Fig. 1).

Reattach the belt clip, if required.
 Attach the connecting cable to the electrodes (Fig. 2).
 The electrodes have a clip fastener to ensure easy connection.





- 7. Place the cable plugs into the socket on the top of the device (Fig. 3).
- 8. Do not pull, twist or make any sharp kinks in the cables (Fig. 4).
- Please note that when the battery is replaced or removed, (i) all settings are restored to the factory default settings.





Fig. 4

6. Program overview

6.1 Basic information

The Digital EMS/TENS unit has a total 50 programs:

- 10 TENS programs
- 30 EMS programs 10 MASSAGE programs

In all programs (see pages 9 to 10) you can set the impulse intensity of both channels individually.

You can also set various parameters in the TENS programs 8-10 and the EMS programs 28-30 to adjust the stimulating effect to the application area.

6.2 Information on placing the electrodes

For the desired success of any stimulation application, it is important that electrodes are placed sensibly.

We advise you to consult with your physiotherapist regarding the best electrode

positions for your intended field of application.

The electrode positions suggested inside the cover (Figs. 1–28) serve as a guide. The following applies when choosing electrode positions:

Electrode distance

The greater the electrode distance which is selected, the greater will be the volume of tissue that is stimulated. This applies to the area and the depth of the tissue volume. At the same time, however, the strength of tissue stimulation decreases as the electrode distance becomes greater, which means that, if a greater electrode distance is chosen, a greater volume is stimulated, but stimulation is weaker. To increase stimulation, pulse intensity must then be

The following is a guideline when selecting electrode distances:

most reasonable distance: approx. 5-15 cm,

below 5 cm, it is mainly surface structures which are strongly stimulated,

over 15 cm, large-area and deep structures are stimulated very weakly. Electrode relationship to muscle fibre orientation Choice of current flow direction must be adapted to the

muscle fibre orientation according to the desired muscle layer. If surface muscles are to be reached, the electrodes must be placed parallel to the fibre orientation (A-B/C-D). If deep tissue layers are to be reached, the electrodes must arrangement can, for example, be achieved via the cross

electrode arrangement, e.g. A-D/B-C. Note the colour-codes of the cables and the channels. The white cable belongs to channel CH1 and the grey cable to channel CH2.

be placed crosswise to the fibre orientation. The latter

In the treatment of pain (TENS) using the Digital EMS/TENS machine with its 2 separately controllable channels and in each case 2 adhesive electrodes, it is advisable to either position the electrodes of one channel so that the pain spot lies between the electrodes or else you position one electrode directly on the pain spot and the other at least 2–3 cm away.

The electrodes of the second channel can be used to treat other pain spots simultaneously or else, however, together with the electrodes of the first channel, to surround the pain area (opposite). A cross arrangement is again advisable here.

Massage function tip: always use all 4 electrodes to ensure optimal treatment.

In order to extend the durability of the electrodes, use them on clean skin which is as free as possible from hair and grease. If necessary, clean the skin with water before application and remove the hair.

If an electrode should come loose during use, the impulse intensity of the corresponding channel is reduced to the lowest level. Apply the electrode again and reset the desired impulse intensity.



6.3 TENS program table

No.	Applications	Phase	(minutes)
-1	Pain relief – acute pain	1	30
2	Pain relief – chronic pain	1	30
	(pulse width modulated)		
3	Endorphin effect (burst)	1	30
4	Sciatica	1	30
5	Treatment of muscle wastage from injury	1	05
		2	15
6	Lumbago	1	20
		2	20
7	Periarthritis	1	15
		3	10

The TENS 8 - 10 program can be set individually, see 7.3.2 (p. 11).

Note: Electrode position is supposed to surround the painful area. With painful muscle groups, the electrodes are grouped round the muscle affected. In case of joint pain, the joint on the front/rear side of the joint and, if electrode distances permit, on the right and left sides of the joint, should be surrounded with electrodes.

Minimal electrode distance should not be less than 5 cm and not exceed 15 cm. See figure no. 25 and 28 for knee and ankle joint.

Burst programs are suitable for all sites which should be treated with an alternating signal pattern (to ensure minimum habituation).

6.4 EMS program table

Applications Applications Possible electrode position Capillarisation Capillarisation 1–28 Warming up Cooling down after training/match Maximum power of lower limbs 22, 23, 24, 26, 27	n n after training/match ower of lower limbs	3	2	1	Prog. No.
Possible electrode position 1–28 1–28 1–28 22, 23, 24, 26, 27		Cooling down after training/match	Warming up	Capillarisation	Applications
	3 2 1 1 1 Phase	1-28	1-28	1-28	Possible electrode position

15 16 17 17								13		n. 12		11	10			9	_		00			7			6			5		No.	
Strengthening of stomach muscles	Strengthening of stomach muscles			Toning of stomach muscles	Lipolysis of stomach muscles	Increasing size of lower limbs		Shaping of lower limbs		Strengthening of lower limbs		Toning of lower limbs	Lipolysis of lower limbs		limbs	Explosive power of torso and upper			Resistance of torso and upper limbs		upper limbs	Maximum power of torso and			Explosive power of lower limbs			Resistance of lower limbs		Chouse and the control of the contro	Applications
10,10,10	10,10,10	18 19 20		18,19, 20	18,19, 20	22, 23, 24, 26, 27	26, 27	22, 23, 24,	26, 27	22, 23, 24,	26, 27	22, 23, 24,	22, 23, 24, 26, 27			1-20			1-20			1-20		26, 27	22, 23, 24,		26, 27	22, 23, 24,	position	electrode	Possible
2		1	2	1	1		2	1	2	1	2	_		ဒ	2	1	ယ	2	-	3	2	1	3	2	-1	3	2	1		1 11000	Phase
0	45	20	10	20	40	15	20	20	10	20	10	20	40	05	12	05	05	12	05	05	15	05	05	12	05	05	20	05	(minutes)	time	Running

Prog. No.	19		20		21		22	23		24		25		26		27	
Applications	Toning of upper limbs		Strengthening of upper limbs		Shaping of upper limbs		Increasing size of upper limbs	Toning of hips and thighs		Strengthening of hips and thighs		Toning of gluteal muscles		Strengthening of gluteal muscles		Shaping of gluteal muscles	
Possible electrode position	12-17		12-17		12-17		12-17	20, 23, 24	ı	20, 23, 24		22		22		22	
Phase	1	2	1	2	1	2	1	1	2	1	2	1	2	1	2	1	2
Running time (minutes)	20	10	20	10	20	20	15	20	10	20	10	20	10	20	10	20	20

 \odot The EMS 28 - 30 programme can be set individually, see 7.3.3 (p. 11).

6.5 MASSAGE program table

Prog.	Applications	Phase	Running time (minutes)
_	Muscle strain relief	1	20
2	Relaxing massage	1	25
ω	Invigorating massage	1	20
4	Tapping massage	1	15
5	Tapping and jet massage	1	15
6	Vibrating massage	1	20
7	Warm up massage 1	1	20
8	Warm up massage 2	1	10
9	Pressure massage	1	20
10	Kneading and pressure massage	•	20

Electrodes must be placed so that they surround the muscle segments in question. For optimal effect, electrode distance must not be greater than approx. 15 cm.

The electrodes should not be applied to the anterior wall of the thorax, i.e. never massage the left or right large pectoral muscle.

Application

7.1 Advice on application

If the machine is not used within 2 minutes, it switches off automatically (automatic switch-off function). When the unit is switched on again, the LCD screen displays the menu selection and the most recently used menu flashes. If an admissible button is pressed, there is one short beep. If an inadmissible button is pressed, there are two short beeps. You can pause the stimulation at any time by briefly pressing the ON/OFF button Ω . To continue the stimulation, briefly press the ON/OFF button Ω again and set the desired impulse intensity again.

7.2 Procedure for programs TENS 1-7, EMS 1-27 and MASSAGE 1-10 (quick start)

Select a suitable program from the program tables (p. 9–10). Place the electrodes on the desired area for treatment (for positioning suggestions see electrode positions, p. 2) and connect them to the device.

- Selecting a program

 Press the ON/OFF button ① to switch on the device.

 Using the M button, navigate through the menus TENS

 MASSAGE (Fig. 1, Example TENS display) and confirm your selection with the E button.

 Using the UP/DOWN buttons, select the desired program and confirm your selection with the E button (Fig. 2, Example TENS program 03).
- If you wish to return to the previous selection menu, press the **M** button. By holding down the **E** button, you can skip individual setting steps and start directly with the stimulation treatment. (i) General information





Fig. 2

Setting the impulse intensity

At the start of stimulation treatment, the impulse intensity of Ch1 and Ch2 is set to 00 as default. No impulses are sent to the electrodes yet.

Using the INTENSITY SETTING buttons, select the desired impulse intensity. The impulse intensity display is adjusted accordingly (Fig. 3). The impulse intensity of Ch1 and Ch2 can be set individually



If the program is in a pause phase, the intensity cannot be increased.
If you wish to end the stimulation treatment early, press the ON/OFF button ①.

7.3 Setting individual parameters

7.3.1 Setting the treatment time

(TENS programs 8-10 and EMS programs 28-30 only) Using the UP/DOWN buttons, select the desired treatment time. You can set a treatment time from 5 minutes to 100 minutes (Fig. 4, Example treatment time of 20 min.). Confirm your selection with the E button. The stimulation treatment begins and the operation status display starts to flash alternately (Fig. 5).

7.3.2 Setting individual TENS programs

Procedure for TENS 08 program

TENS 08 is a preset program that you can also personalise. In this program you can set the impulse frequency to between 1 and 150 Hz and the impulse width to between 50 and 250 µs.

· Place the electrodes on the desired area for treatment (for positioning suggestions see electrode positions, p. 2) and connect them to the device. Select the TENS 08 program, as described in section 7.2 "Selecting a program" (p. 10).

 Using the UP/DOWN buttons, select the desired impulse frequency and confirm with the **E** button (Fig. 1, Example impulse frequency of 100 Hz).

Using the UP/DOWN buttons, select the desired impulse width and confirm with the E button (Fig. 2, Example impulse width of 200 µs).

Select the desired treatment time, as described in 7.3.1 "Setting the treatment time" (p. 11). Select the desired impulse intensity, as described in

7.2 "Setting the impulse intensity" (p. 10).



Fig. 4



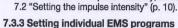
(p. 10). Using the UP/DOWN buttons, select the desired impulse frequency and confirm with the E button (Fig. 1, Example

impulse frequency of 100 Hz). Select the desired treatment time, as described in 7.3.1 "Setting the treatment time" (p. 11).

Select the desired impulse intensity, as described in

 Place the electrodes on the desired area for treatment (for positioning suggestions see electrode positions, p. 2) and connect them to the device.
 Select the TENS 10 program, as described in 7.2 "Selecting a program" 100

Fig. 1



Procedure for the EMS 28 program

Procedure for the TENS 09 program

Procedure for the TENS 10 program

TENS 09 is a preset burst program that you can also personalise. In this program, you can set the impulse width to between 50 and 250 µs.

Place the electrodes on the desired area for treatment (for positioning

and confirm with the E button (Fig. 1, Example impulse width of

TENS 10 is a preset program that you can also personalise. In this program,

you can set the impulse frequency to between 1 and 150 Hz. The impulse width

Select the desired treatment time, as described in 7.3.1 "Setting the treatment time" (p. 11).

Select the desired impulse intensity, as described in 7.2 "Setting the impulse intensity" (p. 10).

changes automatically during the stimulation treatment.

suggestions see electrode positions, p. 2) and connect them to the device. Select the TENS 09 program, as described in 7.2 "Selecting a program" (p. 10). Using the **UP/DOWN** buttons, select the desired impulse width

EMS 28 is a preset program that you can also personalise. In this program, you can set the impulse frequency to between 1 and 100 Hz and the impulse width to between 50 and 320 µs.

Place the electrodes on the desired area for treatment (for positioning suggestions see electrode positions, p. 2) and connect them to the device. Select the EMS 28 program, as described in 7.2 "Selecting a program" (p. 10).





Fig. 2

 Using the UP/DOWN buttons, select the desired impulse frequency and confirm with the E button (Fig. 1, Example

impulse frequency of 30 Hz).

• Using the UP/DOWN buttons, select the desired impulse width and confirm with the E button (Fig. 2, Example impulse width of 250 µs).

Select the desired treatment time, as described in 7.3.1 "Setting the treatment time" (p. 11).

 Select the desired impulse intensity, as described in 7.2 "Setting the impulse intensity" (p. 10).

Procedure for the EMS 29 program

EMS 29 is a preset program that you can also personalise. In this program, you can set the impulse frequency to between 1 and 100 Hz. The impulse width changes automatically during the stimulation treatment.

· Place the electrodes on the desired area for treatment (for positioning suggestions see electrode positions, p. 2) and connect them to the device. Select the EMS 29 program, as described in 7.2 "Selecting a program" (p. 10).

· Using the UP/DOWN buttons, select the desired impulse frequency and confirm with the E button (Fig. 1, Example impulse frequency of 30 Hz).

Select the desired treatment time, as described in 7.3.1 "Setting the treatment time" (p. 11).

Select the desired impulse intensity, as described in 7.2 "Setting the impulse intensity" (p. 10).

Procedure for the EMS 30 program

EMS 30 is a preset program that you can also personalise. In this program, you can set the impulse frequency to between 1 and 100 Hz. You can also set the working time and pause time for this program to between 1 and 30 seconds

 Place the electrodes on the desired area for treatment (for positioning suggestions see electrode positions, p. 2) and connect them to the device. Select the EMS 30 program, as described in 7.2 "Selecting a program"



Fig. 1



Fig. 2

Fig. 1

Using the **UP/DOWN** buttons, select the desired length of pause time and confirm with the **E** button (Fig. 2, Example

Using the UP/DOWN buttons, select the desired length of

working time and confirm with the E button (Fig. 1, Example

pause time of 10 seconds).
Using the UP/DOWN buttons, select the desired impulse frequency and confirm with the E button (Fig. 3, Example impulse frequency of 30 Hz).

Select the desired treatment time, as described in 7.3.1 "Setting the treatment time" (p. 11).

Select the desired impulse intensity, as described in 7.2 "Setting the impulse intensity" (p. 10).





Fig. 3

7.4 Changes to the settings

working time of 2 seconds).

Changing intensity (during application)

• Cit +/- und Cit +/-: Changing the intensity for each channel.

Interrupting stimulation Press the ON/OFF button ①.

Changing application (completely or single parameters) ON/OFF button ①: to interrupt stimulation

Setting the program, see 7.2; setting individual parameters, see 7.3.

7.5 Doctor's Function

Doctor's Function is a specific setting to allow you to call up your specific personal program even more easily.

Your individual program settings are instantly recalled and activated when the device is switched on.

Settings for this individual program may, for example, be on the advice of your doctor.

Setting the Doctor's Function

- Select your program and appropriate settings as described under 7.2 or 7.3.
- At the start of the stimulation treatment, the impulse intensity of Ch1 and Ch2 is set to 00 as default. No impulses are sent to the electrodes yet.

Before setting the desired impulse intensity using the intensity setting buttons, press and hold the Ch2 - button for 5 seconds. Storage in the Doctor's Function is confirmed with a long acoustic signal.

Cancelling Doctor's Function

To clear the device again and to re-allow access to other programs, press and hold the Ch2 - button again for approx. 5 seconds. To do this, the impulse intensity of Ch1 and Ch2 must be set to 00. Deletion of the Doctor's Function is confirmed with a long acoustic signal.

8. Cleaning and storage

Adhesive electrodes

- To ensure that the adhesive electrodes remain adhesive for as long as possible, clean them carefully with a damp, lint-free cloth or clean the underside of the electrodes under lukewarm running water and pat dry with a lint-free cloth.
 - Before cleaning with water, remove the connection cables from the
- · After use, stick the electrodes onto the backing film.

Cleaning the unit

- Remove the batteries from the device every time you clean.
- · Clean the unit after use with a soft, slightly moistened cloth. In case of more extreme soiling you can also moisten the cloth with mild soapy water.
- Do not use any chemical cleaners or abrasive agents for cleaning.

_____ Ensure that no water enters the device.

Storage

- · Remove the batteries from the unit if you are not going to use it for a longer period. Leaking batteries can damage the unit.
- Do not make any sharp kinks in the connecting leads or electrodes.
- Detach the connecting cable from the electrodes.
- · After use, stick the electrodes onto the backing film.
- Store the machine in a cool, well-ventilated place.
- Never place any heavy objects on the machine.

9. Disposal

Used, fully discharged batteries must be disposed of in a specially labeled collection container, at toxic waste collection points or through an electrical retailer. You are under legal obligation to dispose of batteries correctly. Note: You will find these markings on batteries containing harmful substances: Pb = battery containing lead, Cd = battery containing

cadmium, Hg = battery containing mercury.
Please dispose of the device in accordance with the directive 2002/96/EC - WEEE (Waste Electrical and Electronic Equipment). If you have any queries, please refer to the local authorities responsible for waste disposal.



10. Problems and solutions

The machine does not switch on when the ON/OFF button ① is pressed. What should I do?

- (1) Make sure that the batteries have been inserted correctly and have contact.
- (2) Change the batteries if necessary. (3) Contact customer service.

The electrodes are detaching themselves from the body. What should I do?

- (1) Clean the adhesive surface of the electrodes with a damp, lint-free cloth. Then let them dry in the air and attach them again. Should the electrodes still not adhere firmly, they must be replaced.
- (2) Before each application, clean the skin, avoiding using any skin cream or oils before treatment. Shaving can increase adhesion of the electrodes.

There is no noticeable stimulation. What should I do?

- (1) Interrupt the program by pressing the ON/OFF button ①. Make sure that the connecting leads are properly connected to the electrodes. Make sure that there is a firm contact between the electrodes and the treatment area.
- (2) Make sure that the plug of the connecting lead is firmly connected to the machine.
- (3) Press the ON/OFF key 10 to start the program again.
- Check that the electrodes are positioned properly and make sure that the adhesive electrodes are not overlapping.
- (5) Increase pulse intensity in stages
- (6) The batteries are almost empty. Replace these.

What to do if the battery symbol is displayed.

Replace all batteries.

You are aware that the electrodes feel uncomfortable. What should I do?

(1) The electrodes are badly positioned. Check positioning and if necessary reposition the electrodes.



(2) The electrodes are worn. Due to the fact that current distribution can no longer be guaranteed evenly over the entire surface, these may lead to skin irritation. You must therefore replace these.

The skin in the treatment area becomes red. What should I do?

Stop treatment immediately and wait until the skin has returned to normal. If any redness beneath the electrode soon disappears, this is not dangerous and is due to the increased circulation which has been stimulated locally. If, however, the skin irritation remains and there is possibly itching or inflammation, please consult your doctor before further use. It may possibly be caused by an allergy to the adhesive surface.

11. Technical details

TENS Digital Pain Relief unit Name:

Model: EM43

Initial curve shape: biphase square-wave pulse

Pulse duration: 50-450 µs Pulse frequency:

1–150 Hz max. 100 Vpp (at 500 Ohm) max. 200 mApp (at 500 Ohm) 3x AAA batteries Output voltage: Output current:

Power supply:

Treatment time: adjustable from 5 to 100 minutes

Intensity: adjustable from 0 to 50

Operating conditions: 10°C-40°C (50°F-104°F) with a relative humidity

of 30–85 % $-10\,^{\circ}\text{C}-50\,^{\circ}\text{C}$ (14 $^{\circ}\text{F}-122\,^{\circ}\text{F}$) with a relative humidity of 10–95 % Storage conditions:

135 x 66 x 29 mm (incl. belt clip) Dimensions:

107 g, (incl. belt clip, without batteries)
133 g, (incl. belt clip and batteries) Weight:

Explanation of symbols: Application part type BF

Caution! Read the instructions for Use.

Note: If the machine is not used according to these specifications, perfect functioning is not guaranteed.

We reserve the right to make any technical alterations that are necessary in order to improve and develop the product further.

This unit is in line with European Standards EN60601-1, EN60601-1-2

and EN60601-2-10 and is subject to particular precautions with regard to electromagnetic compatibility (EMC). Please note that portable and mobile HF communication systems may interfere with this unit. For more details, please contact your local Boots store.

The machine meets the requirements of the European Medical Device Directive 93/42/FC

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Electromagnetic Compatibility Information

Table 1

Guidance and manufacture's declaration – electromagnetic emissions

The EM 43 is intended for use in the electromagnetic environment specified below.

The customer or the user of the EM 43	should assure that	t it is used in such an environment.
Emissions test	Compliance	Electromagnetic environment - guidance
RF emissions CISPR 11	Group 1	The EM 43 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 emissions CISPR 11	Class B	The EM 43 is suitable for use in all establishments, including domestic establishments and those directly connected to the public
Harmonic emissions IEC 61000-3-2	Not applicable	low-voltage power supply network that supplies buildings used for domestic purposes.
Voltage fluctuations/flicker emissions IEC 61000-3-3	Not applicable	

Guidance and manufacturer's declaration – electromagnetic immunity of the EM 43 should assure that it is used in such an environment.

IMMUNITY test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
			Portable and mobile RF communications equipment should be used no closer to any part of the EM 43, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
			Recommended separation distance:
Conducted RF IEC 61000-4-6	3V _{ms} 150 kHz to 80 MHz	3V	3V $d = 1,2 \sqrt{P}$ 80 MHz to 800 MHz $d = 2,3 \sqrt{P}$ 800 MHz to 2,5 GHz
Radiated RF IEC 61000-4-3	3V/m 80 MHz to	3V/m	Where <i>P</i> is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and <i>d</i> is the recommended separation distance in meters (m).
	2,5 GHz		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. 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.

* 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 EM 43 is used exceededs the applicable RF compliance level above, the EM 43 should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the EM 43.

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

Table

The FM 43 is intended for use in the e	lectromagnetic environment so	pecified below. The customer of	or the user of the EM 43 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	± 1 kV for input/output lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	± 1 kV line to line ± 2 kV line to earth	Not applicable	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_{τ} (>95% dip in U_{τ}) for 0.5 cycle 40% U_{τ} (60% dip in U_{τ}) for 5 cycles 70% U_{τ} (30% dip in U_{τ}) for 25 cycles <5% U_{τ} (>95% dip in U_{τ}) for 15 cycles	Not applicable	Mains power quality should be that of a typical commercial or hospital environment. If the user of the EM 43 requires continued operation during power mains interruptions, it is recommended that the EM 43 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.

Table 4

Recommended separation distances between portable and mobile RF communications equipment and the EM 43

The EM 43 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the EM 43 can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the EM 43 as recommended below, according to the maximum output power of the communications equipment.

		Separation distance according to fr	equency of transmitter	
Rated maximum output power of transmitter (W)	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	00	

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be determined 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.