USER MANUAL

Shockwave Therapy Device SW3200

ShockPhysio Pro

IFU-SW3200-NULL-V1.5-EN

CE 0598

EN

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1 Foreword

This manual is valid for the Shenzhen Dongdixin Technology Co., Ltd. (Hereinafter referred to as Dongdixin) Shockwave Therapy Device.

Welcome to using our Shockwave Therapy Device!

This user manual provides comprehensive information on the device's performance indicators, usage guidelines, and maintenance procedures. It is an essential document that should be kept close to the device for reference. Before installing and utilizing the Therapy Device, the user of this device must read this user manual carefully to ensure proper usage, achieve optimal performance, and comply with prescribed safety standards.

Please store this instruction manual in a convenient place or keep it alongside the device for future reference.

The specifications mentioned in this manual are accurate at the time of publication. However, as part of our commitment to continuous improvement, Dongdixin reserves the right to make changes to these specifications without prior notice.

FCC Regulatory Compliance

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

This equipment has been tested and found to comply with the limits for a Class B digital device, pursuant to Part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation.

If this equipment does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:
-- Reorient or relocate the receiving antenna.
-- Increase the separation between the equipment and receiver.
-- Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.
-- Consult the dealer or an experienced radio/TV technician for help.

RF Exposure Compliance(Main unit)

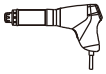
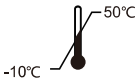
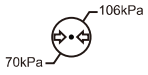
This equipment complies with FCC radiation exposure limits set forth for an uncontrolled environment. This equipment should be installed and operated with minimum distance 20 cm between the radiator and your body. This transmitter must not be co-located or operating in conjunction with any other antenna or transmitter.

RF Exposure Compliance(Wireless decive)

The device has been evaluated to meet general RF exposure requirement. The device can be used in portable exposure condition without restriction.
This device and its antenna(s) must not be co-located or operation in conjunction with any other antenna or transmitter.

1.1 Symbols and Markings

The markings on the device are assurance of its conformity to the highest applicable standards of medical equipment safety and electromagnetic compatibility and conform to ISO 7010 and ISO 15223-1. One or more of the following markings may appear on the device:

Symbols	Meaning
	Wired Handpiece
	Stand-by
O/I	Power OFF/ON
	Type BF applied part
	Refer to instruction manual
	Operating instructions
	Keep dry
	Stacking limit by number
	Transportation and storage temperature limit
	Transportation and storage humidity limitation
	Transportation and storage atmospheric pressure limitation
	General symbol for recovery/recyclable
	Manufacturer

	Date of manufacture
CE 0598	CE mark
EC REP	Authorized representative in the European Community
SN	Serial number
LOT	Batch code
MD	Medical device
UDI	Unique device identifier
	Disposal in accordance with Directive 2012/19/EU (WEEE)
	Caution
CE	Marked in the Battery means that the battery complies with the European Regulation (EU) 2023/1542. Marked in the component means that the component complies with the corresponding European Regulation (EU).
REF	Catalogue number or reorder number or part number
	Fragile, handle with care
	This way up
	The device is MR Unsafe
	Charging holder

2 Safety and Indications

2.1 Intended Use

2.1.1 Indications

The Shockwave Therapy Device is a treatment system for the electromagnetic generation and application of radial shock waves in orthopaedics and physiotherapy.

Indications: The Shockwave Therapy Device is used to relieve pain and provide treatment of myofascial pain syndrome, plantar fasciitis and patellar tendinopathy.

2.1.2 Contraindications

The device should not be used:

- Vascular diseases present in or near the treatment area.
- Local infections in the treatment area.
- Around malignant or benign tumours.
- Directly on cartilage surfaces or near the small facet joints of the spinal column.
- Directly over implanted electronic devices such as pacemakers, analgesic pumps etc.
- In areas, in which mechanical energy in the form of vibrations may lead to tissue damage such as metal implants after a fracture.
- Open wounds in or near the treatment area.
- After fractures, torn muscle fibres or muscle tears.

In general we advise against treatment:

- If blood clotting disorders are present or the patient is receiving treatment that results in a change in the blood clotting behaviour.
- During pregnancy.
- On patients with neurological diseases resulting in impairment of the vasomotor function in the treatment area.
- Over air-filled cavities such as treatment on the thoracic spine, etc.
- On children, particularly around the epiphyseal plates.

Care is required for patients:

- With impaired sensibility.
- With severe autonomic disorders.
- Under the influence of drugs and/or alcohol as circulatory stresses and inadequate treatment responses cannot be excluded.

2.1.3 Risks and Side Effects

Side effects could occur after a treatment with Radial Pressure Wave therapy.
The majority will appear after 1-2 days. Do not repeat a treatment until the previous side effects have diminished. Common side effects include:

- Erythema, reddening and discomfort.
- Swelling.
- Pain.
- Hematoma.
- Petechiae, red spots.
- Irritation.

Skin lesions after previous cortisone therapy.
These side effects generally abate after 5 to 10 days.

Residual Risk

For this device, the noise level of the shockwave pulse is within safe area. The users and patients may feel upset after a period of treatment.

2.2 Intended User Profile

Shockwave Therapy Device is intended exclusively for use by licensed or qualified medical specialists and is only allowed to be used by qualified and trained medical persons.
Operators of the shockwave Therapy Device must have been adequately trained in using this device safely and efficiently before they operate the device described. An introduction to the principles of operation is provided by the Dongdixin dealer with reference to the user manual. The operator must read this manual before operating the device to understand its use principle, contraindications and side effects, etc.

2.3 Intended Patient Profile

The intended patients to be treated are patients between 18 and 65 years old, suffering from muscular soreness, myofascial pain syndrome, plantar fasciitis and patellar tendinopathy.

2.4 Intended Environment for Use

The device is designed for operation in a clinical setting and can be moved between rooms. It is not intended for frequent transport between different facilities.

2.5 Precautionary Instructions

In this section general Precautions and Warnings are listed, that you should be aware of when using the device.
The definition of these symbols including danger, warning, caution and NOTE are as follows:

⚠ WARNING

Refer to a situation of potential danger which, if not avoided, could lead to serious injury.

⚠ CAUTION

Indicate that incorrect operation could lead to minor injuries.

NOTE:
Refer to the additional information concerning specific features or operating instructions.

⚠ WARNING

- This device should only be used by licensed or qualified medical specialists.
- This device is designed to only be used by trained medical professionals who use the medical device in the course of their work and in the framework of a professional healthcare activity, and understand the benefits and limitations of shockwave therapy. For example, physical therapist, occupational therapist, sport medicine therapists, medical doctors and so on.
- This device should only be used under the continued supervision of a physician or licensed practitioner.
- Users of this device must be trained in how to use the system properly and have the appropriate skills.
- Any treatment instructions regarding treatment location, duration and strength require medical knowledge and should only be given by authorized doctors, therapists and health paraprofessionals. It is imperative that these instructions are followed.
- Treatment must always be carried out under medical supervision.
- The device is not designed for persistent use. After a treatment with maximum 6000 shocks, a break of 15 minutes becomes necessary.
- The device is not designed to prevent the ingress of water or liquids. Ingress of water or liquids could cause malfunction of internal components of the system and therefore create a risk of injury to the patient.
- To prevent the risk of electric shock, ensure that the device is electrically grounded by connecting it to a grounded electrical service receptacle in accordance with the applicable national and local electrical codes.

- Operation of this equipment in the vicinity of strong electromagnetic fields (e.g. tomography, x-ray or diathermy equipment) may interfere with the operation of the device. Please keep a safe distance of several meters.
- Modification of this equipment is strictly prohibited. Do not make any changes to this equipment without authorization from the manufacturer.
- Exercise caution when operating this equipment around other devices. Potential electromagnetic or other interference could occur with this equipment or other devices. Minimize this interference by avoiding the simultaneous use of other equipment.
- The instruments must only be operated with the mains power cable provided. Protect the mains cable from any mechanical stress.
- This equipment is not suitable for use in the presence of flammable anesthetics mixed with air, oxygen, or nitrous oxide. There is a risk of an explosion.
- It must not be used in wet areas. If it is used in wet areas, significant damage may occur, and patients and users may be endangered.
- Do not attempt to service or maintain the device while it is in use.
- Please remove the battery if the device is not likely to be used for some time.

⚠ CAUTION

Please read, understand, and follow the precautionary and operating instructions carefully. Familiarize yourself with the limitations and potential hazards associated with using any Radial pressure wave device. Pay attention to the precautionary and operational decals placed on the device.

- Operate this device within the temperature range of 5°C to 30°C (41°F to 86°F) and maintain a Relative Humidity between 20% and 80%.
- Transport and store this device at -10°C to 50°C (14°F to 122°F) and between 10% and 93% Relative Humidity.
- Ambient pressure: 700 hPa to 1060 hPa.
- Avoid exposing the device to direct sunlight, heat from radiators, excessive dust, moisture, vibrations, and mechanical shocks.
- If any liquid enters the device, disconnect it from the mains supply immediately and have it checked by authorized people.
- Before administering any treatment to a patient, thoroughly understand the operating procedures for each treatment mode, as well as the indications, contraindications, warnings, and precautions. Seek additional information from reliable sources on the application of Radial pressure wave therapy.

- If you encounter any issues with this device, such as setup, maintenance, or operation, please contact the manufacturer or dealer for assistance. Additionally, please report any unexpected operation or events to your manufacturer or dealer.
- Treatment with device may occasionally cause irritation, petechiae, bruising, swelling, pain, discomfort and redness on the treatment area.
- Caretakers should assess patients for adverse reactions on the skin where the device has contact, such as redness (erythema), swelling (edema), irritation, sensitization (delayed Type IV hypersensitivity), allergy, immune response, or other reactions.
- You may feel upset by projectile noise after using the applicator for a long time.

NOTE:
The SW3200 may only be used with the intended components and accessories, i.e. the wired handpiece and battery pack.

3 Product Description

The Shockwave Therapy Device SW3200 is an electromagnetic operated ballistic shock wave generator. Accelerated by electromagnetic, the movement of projectile in the precision ballistic mechanism of handpiece generate the shockwave. This shockwave will be transferred to the tissue with the help of the gel, and then conduct therapeutic effect on the tissue.

3.1 Scope of Supply

Part No.	Description
SW3200.2-IN-1	Shockwave Therapy Device SW3200 2 in 1
SW3200.4-IN-1	Shockwave Therapy Device SW3200 4 in 1

Standard Accessories for SW3200 2 in 1:

Part No.	Description
AC9017.200	Wired Handpiece
AC9018.200	Wireless Handpiece
AC9103.300	Applicator SR15 x2
AC9106.300	Applicator CR15 x2
AC9107.300	Applicator CR20 x2
ACC050C706	Charging Holder
ACC050C701	Wired Handpiece Holder
330050000012	Charging Cable
ACC050C600	Rechargeable Battery
ACC050C707	Working Platform
ACC060C300	Applicator Removal Tool
ACC010B300	Power Cable
ACC020C800	O-ring(φ 9.5-φ 13.5mm) x 10
ACC020C801	O-ring(φ 15-φ 19mm) x 10
ACC020C802	O-ring(φ 20-φ 24mm) x 10
/	PC Sticker x2
SW3200B100	User Manual (USB Driver)

Standard Accessories for SW3200 4 in 1:

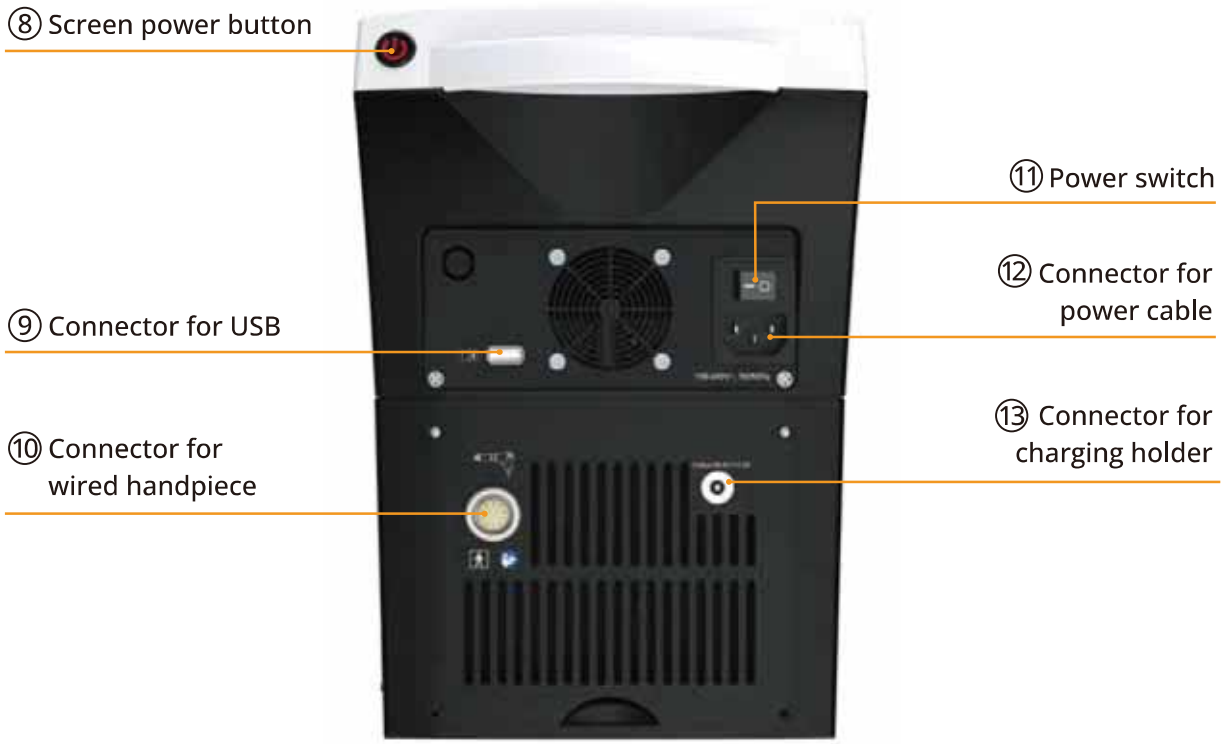
Part No.	Description
AC9017.200	Wired Handpiece
AC9018.200	Wireless Handpiece x3
AC9103.300	Applicator SR15 x4
AC9106.300	Applicator CR15 x4
AC9107.300	Applicator CR20 x4
ACC050C706	Charging Holder
ACC050C701	Wired Handpiece Holder
330050000012	Charging Cable
ACC050C600	Rechargeable Battery x3
ACC050C707	Working Platform
ACC060C300	Applicator Removal Tool
ACC010B300	Power Cable
ACC020C800	O-ring(φ 9.5-φ 13.5mm) x 10
ACC020C801	O-ring(φ 15-φ 19mm) x 10
ACC020C802	O-ring(φ 20-φ 24mm) x 10
/	PC Sticker x2
SW3200B100	User Manual (USB Driver)
CART200.301	CartRehab

Optional Accessories:

Part No.	Description
AC9100.300	Applicator CD15
AC9101.300	Applicator CD20
AC9102.300	Applicator CD35
AC9104.300	Applicator SR20
AC9105.300	Applicator SR35
AC9108.300	Applicator CR35
ACC020C803	O-ring(φ 35-φ 39mm)
CART200.301	CartRehab

3.2 Operator Interface

SW3200 2 in 1



SW3200 4 in 1


① LCD touchable display
② Wireless Handpiece
③ Charging Holder
④ Adjustment knob
⑤ Wired Handpiece
⑥ Wired Handpiece Holder
⑦ Working Platform


⑧ Screen power button
⑫ Power switch
⑨ Connector for USB
⑬ Connector for power cable
⑩ Connector for wired handpiece
⑭ Connector for charging holder
⑪ Trolley

Wired Handpiece

The handpiece contains the shockwave generator, a fan to dissipate heat and the slot for the different applicator. It is connected to the control main device.

Wireless Handpiece

Indicator Light

① CH3: The current wireless handpiece is connecting to channel 3

② CH2: The current wireless handpiece is connecting to channel 2

③ CH1: The current wireless handpiece is connecting to channel 1

④ Ramp up switch light

⑤ RAMP UP: RAMP UP button

⑥ POWER: Power button

Program light

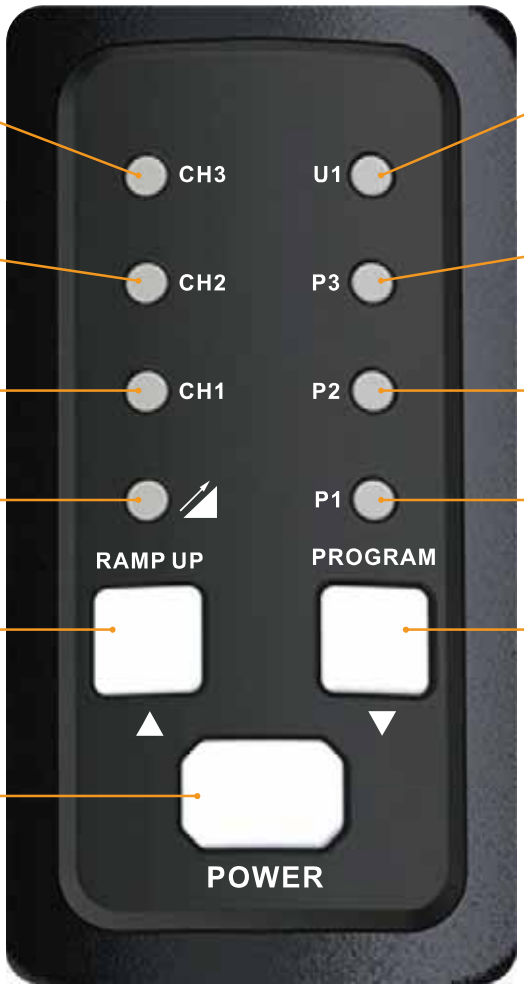
⑦ U1: Custom protocols is downloaded from the main device

⑧ P3: Pre-set program 3

⑨ P2: Pre-set program 2

⑩ P1: Pre-set program 1

⑪ PROGRAM: Program button

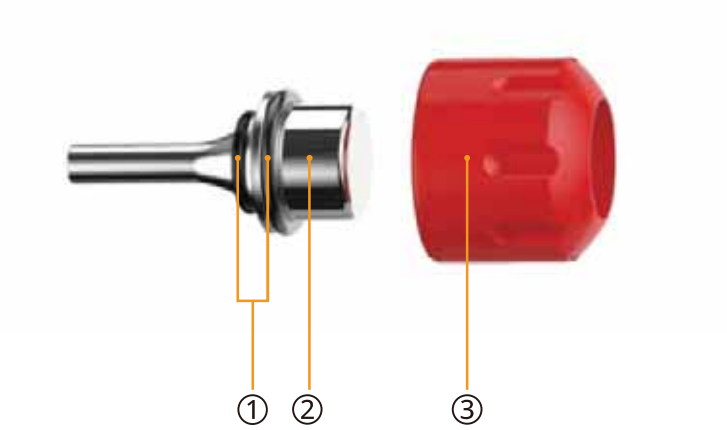


⚠ CAUTION

- When using the handpiece on a patient, it is crucial to securely screw one of the applicator onto the handpiece, ensuring it is tightened as much as possible.
- The cable should not be stretched beyond its maximum length and must be protected from any pinching or mechanical damage during usage.
- To prevent the accumulation of heat in the handpiece, it is essential to avoid obstructing the air vents located at the top and base of the handpiece with the hand or any other object. This will allow proper ventilation and maintain optimal performance.

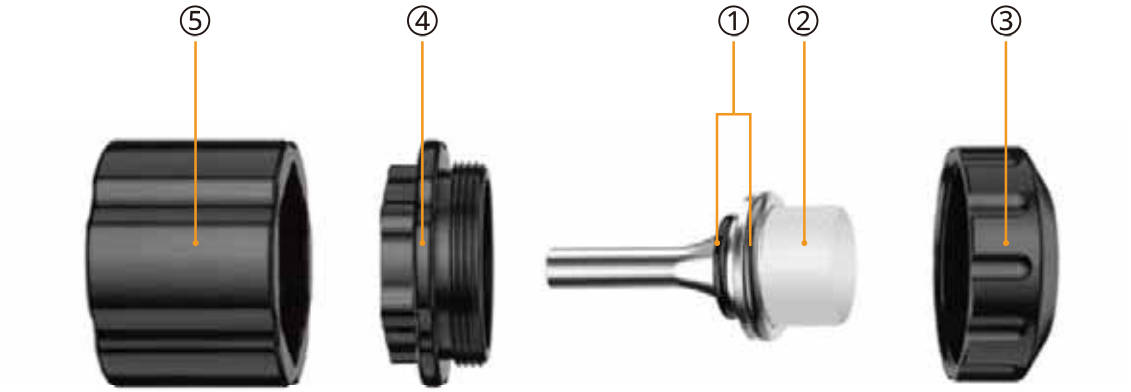
Applicator

Size 15mm



① O-ring
② Applicator insert
③ Applicator Screw Cap
④ Applicator inner Screw Cap
⑤ Applicator Removal Tool

Sizes 20mm and 35mm



NOTE:
Pictures of wired handpiece and applicator are examples. Individual components may be different from those shown in the illustration.

Depending on the therapy to be performed, the handpiece can be equipped with different applicator. Dongdixin has a total of 9 applicators to choose from, please contact with manufacturer or dealer for specific applicator specifications. The difference between treatment heads is as follows:

- SR Series: Steel Round applicator, the contact part of the applicator is composed of steel and the contact surface is rounded. The SR series is available in 3 sizes (15mm, 20mm and 35mm).
- CR Series: Compound Round applicator, the contact part of the applicator is composed of steel and silica gel composite materials and the contact surface is rounded. The CR series is available in 3 sizes (15mm, 20mm and 35mm).
- CD Series: Compound Deep applicator, the contact part of the applicator is composed of steel and silica gel composite materials and the contact surface is flat. The CD series is available in 3 sizes (15mm, 20mm and 35mm). The applicator is used to transmit the shockwaves.


SR 15


SR 20


SR 35


CR 15


CR 20


CR 35


CD 15

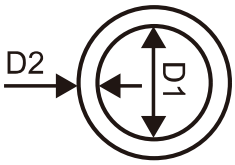

CD 20


CD 35

NOTE:
On the impact rod part of each applicator, there is its corresponding model radium engraving identification.

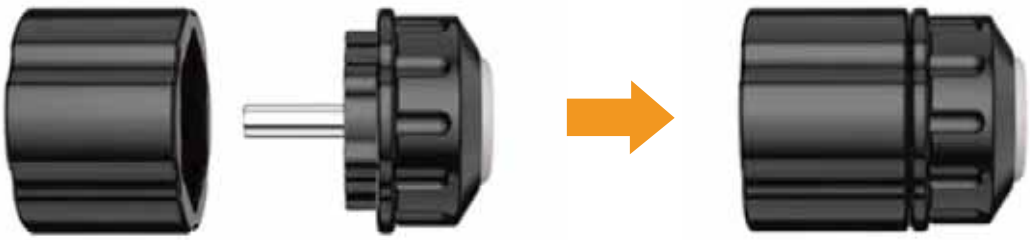
O-ring

The O-ring is used to cushion the applicator while the handpiece is working. Each applicator is equipped with two O-rings, as shown in the diagram below. The O-ring type configured for each applicator is different.

Ref	Part No.	O-ring	Measurements	Used for	
1	ACC020C800	D1=9.5mm D2=2.0mm		Applicator SR15/20/35 Applicator CR15/20/35 Applicator CD15/20/35	
2	ACC020C801	D1=15mm D2=2.0mm		Applicator SR15 Applicator CR15 Applicator CD15	
3	ACC020C802	D1=20mm D2=2.0mm		Applicator SR20 Applicator CR20 Applicator CD20	
4	ACC020C803	D1=35mm D2=2.0mm		Applicator SR35 Applicator CR35 Applicator CD35	

Applicator Removal Tool

The applicator removal tool is used to remove and install the 20/35 mm applicator.



⚠ CAUTION

Connection of accessories other than the ones specified by the manufacturer can adversely affect the safety of the patient and correct functioning of the equipment, and is therefore not permitted.

3.3 Device Light Indicators

4 Installation Instructions

4.1 Transport and Storage Conditions

Ambient temperature: -10°C to +50°C (14°F to 122°F).
Relative humidity: 10% to 93%.
Ambient pressure: 700 hPa to 1060 hPa.

4.2 Remove Packaging

Upon delivery, the device comes packaged with materials provided by the manufacturer. With a weight of approximately 4 kg for main device and a weight of approximately 14 kg for trolley, it can be unpacked by a single person. Follow these steps to unpack the device:

• Position the transport packaging in right way, make sure the arrows are upward.
• Remove the safety bands securing the transport packaging.
• Lift the device inner box out from the transport packaging.
• Open the device inner box.
• Inspect the device for any signs of damage before proceeding.

Complete the following steps immediately after unpacking:
• Check whether the outer packaging box is complete and damaged.
• Verify the delivery document to ensure all items have been received.
• Inspect the external items and accessories for any potential damage from transport.
• Confirm that the packaging includes the complete scope of supply as outlined in Chapter 3.1.
• Retain the original packaging, as it may be useful for transporting the device in the future.

NOTE:
If you find anything is damaged, please contact your dealer or manufacturer immediately.

4.3 Set-up

NOTE:
• Please perform the following steps to install the device.
• Before starting up the system, remove the device from its transport packaging. Do not operate the device while it is in the packaging.
• Make sure that the Power Switch on the device is set to 'O'.

Installation of SW3200 4 in 1

• Take out the main device from the transport case.
• Take out the trolley from the transport case. Install the trolley following the instruction within the package.
• Install the main device to the working platform and trolley following the QUICK ASSEMBLY GUIDE within the main device package.
• Place the trolley with main device on a stable surface.

Installation of SW3200 2 in 1

With CartRehab:
• Take out the main device from the transport case.
• Take out the trolley from the transport case. Install the trolley following the instruction within the package.
• Install the main device to the working platform and trolley following the QUICK ASSEMBLY GUIDE within the main device package.
• Place the trolley with main device on a stable surface.

Without CartRehab:

- Take out the main device from the transport case.
- Install the main device to the working platform following the QUICK ASSEMBLY GUIDE within the main device package.
- Place the main device on a stable surface.

Power Up

1. Insert the power cord into the back of the device, and plug it into the power outlet.
2. Turn on the device using the power switch located at the back.

Power Off

To completely power off the device, turn off the main device power switch and disconnect the power cable from the socket.

NOTE:
Ensure that the power switch remains accessible at all times, as it can function as an emergency switch.

Turn on /off for Wireless handpiece

Turn on: Short-press the POWER button, handpiece turns on and indicator light lightened. If no operation for 3 minutes (30s when low battery), the handpiece will automatically turn off.
Turn off: Long-press the POWER button to turn off.

4.4 Connection of Accessories

Connecting the Handpiece

• Plug the handpiece into the connector for wired handpiece of the device.

Make sure that the red dot on the handpiece plug is aligned with the red dot on the connector of device.

Connect the charging cable

Insert the charging cable into the “Charging Cable Output Interface” on the main device.

Place the wireless handpiece

Refer to Chapter 5.2 to pairing the wireless handpiece before use.

Align the socket of wireless handpiece with the DC plug in the charging holder, then put the wireless handpiece onto the charging holder.

NOTE:

The charging status of wireless handpiece is as follows:

Green light: indicates the wireless handpiece is fully charged.

Red light: indicates the wireless handpiece is charging.

4.5 Replacement of Accessories

Replacement of Applicator

NOTE:

The applicator replacement method for the wired handpiece and wireless handpiece is the same .

Take wired handpiece as example.

To replace the different applicator, please follow these steps:

1. Disconnect the handpiece from the main device, and then hold the handpiece in one hand.
2. With the other hand, unscrew the current applicator from the handpiece in an anticlockwise direction .

3. Take out the desired applicator from the accessory box and securely screw it onto the handpiece in a clockwise direction.

4. Continue tightening until the black outside ring of the applicator rests flush against the handpiece, tighten clockwise until it finger-tight, and pay attention not to use excessive force.
5. Ensure that the applicator is firmly attached to the handpiece to prevent any potential issues during treatment.

NOTE:
There are three different sizes of applicator, there is a small difference when removing and installing the 15mm applicator and when removing and installing the 20/35 mm applicator.

Before installing the 20 or 35 mm applicator, please screw the applicator inner screw cap into the applicator screw cap tightly, shown as below:

① ② ③

⚠ CAUTION

- Please note that applicator are consumable parts and should be replaced after a specific period of use to maintain optimal performance.
- Minor or slight deformation or shortening of the rear impact dome will not impact the functionality of the applicator.
- However, if you notice significant deformation or substantial shortening of the rear impact dome, it is essential to replace the applicator promptly to ensure safe and effective treatments.
- If the applicator has worked more than 300,000 times, it is recommended to check at least once a week.

Replace Rechargeable Battery

The wireless handpiece is powered by a rechargeable battery. To ensure the wireless handpiece always functions optimally, it's necessary to replace the battery when its lifespan ends or when the battery's performance begins to decline. Typically, after approximately 200 discharge cycles, the battery's capacity and discharge capabilities start to decrease.

1. Firstly, ensure that the wireless handpiece is turned off.
2. Locate the battery compartment at the end of the wireless handpiece and use a screwdriver to turn counterclockwise to unscrew the screw of the battery cover.

3. Pinch both sides of the battery cover with your hands and force back to remove the battery cover.

4. Pinch both sides of the battery and force backwards to pull it out.

5. Place the new battery in the battery compartment. Please be sure to align the battery to the slide rail in the battery compartment and take care to correctly align the polarity of the battery.

6. Then push the battery forward. The forward push can be stopped until the inhibiting device of the battery contacts the edge of the battery compartment.

7. Finally, re-tighten the screw of the battery cover to complete the battery replacement (Screw clockwise). Please ensure that the seal is secure.

NOTE:
Disposal of the rechargeable battery and its components must be carried out in accordance with national waste disposal regulations.

5 Operations

5.1 User Interface

③ Screen power button: When the device is working, press this button to stop the work. When the device is in standby state, press this button to hibernate the device.

Device Power On

Power Supply from Power Outlet: Insert the power cable into the power socket of the main device, and then connect the other end to the power outlet. Turn on the power switch, the device powers on.

The Initialization screen below will be shown for a few seconds whilst the device starts.

Startup Picture

Device power off

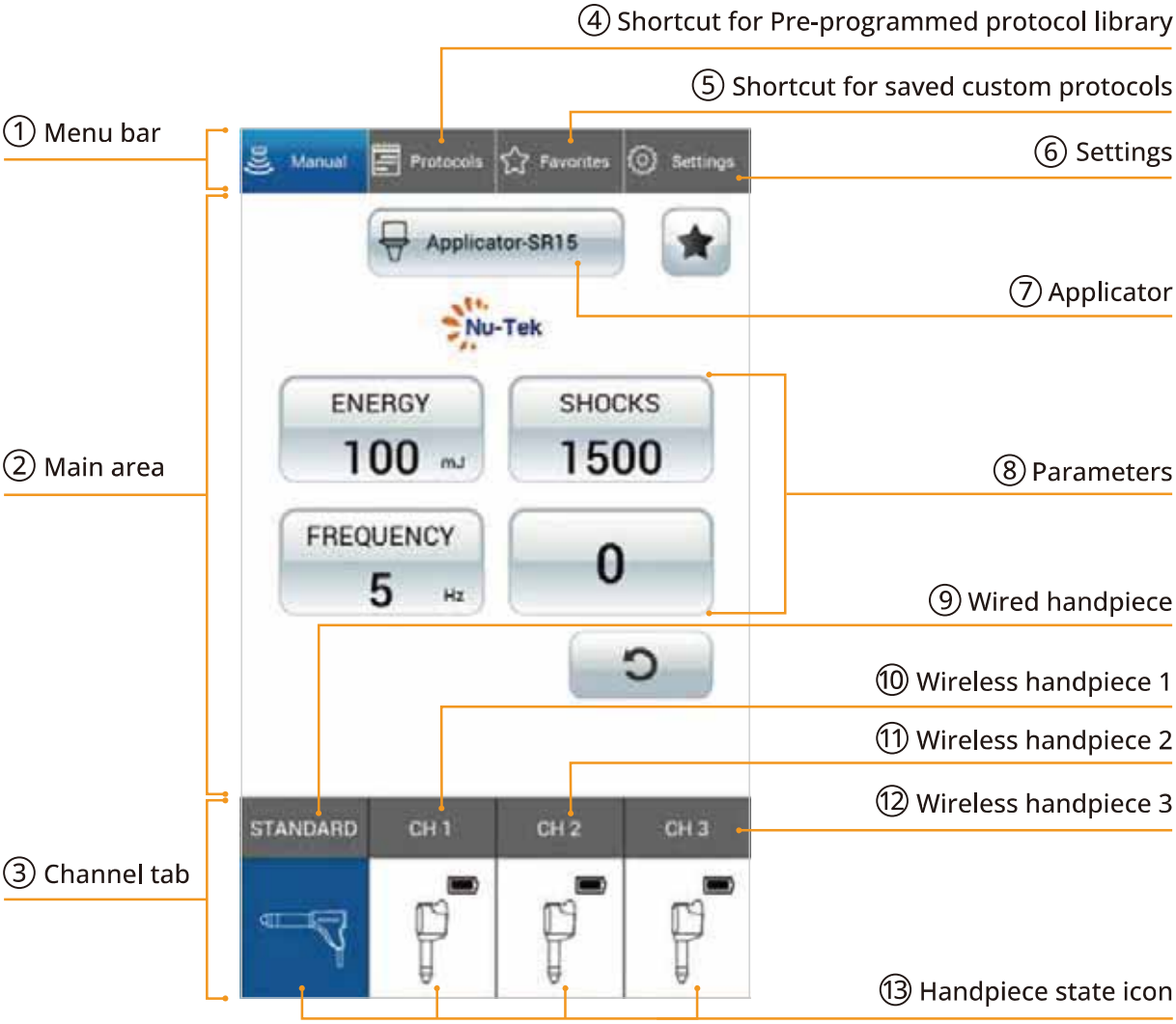
Power Supply from Power Outlet: Turn off the power switch to power off the device, then unplug the power cable.

NOTE:

The following software screens take SW3200 4 in 1 as example. The software screen difference of SW3200 4 in 1 and SW3200 2 in 1 is the number of wireless handpiece icon. SW3200 2 in 1 only has one wired handpiece icon and one wireless handpiece icon.

Manual screen

Manual screen provides access to manual operation of therapy. The therapy screen has the following information.

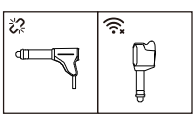


⑦ Applicator: Touch to select the applicator you use

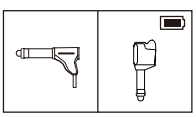
⑧ Parameters: Touch to activate the parameter box, once it become deep blue color, you can adjust it with adjustment knob

⑬ Handpiece state icon:

Disconected: Handpiece not connected



Ready/Working: Handpiece connected or handpiece is working



NOTE:

Only paired handpiece will be displayed on the screen.

5.2 System Settings

Press  to adjust preferences.

Language:

The device supports multiple languages. Click the "Language button" and a language list window will appear. Then select the language you want. The device includes English, Spanish, French, German, Italian, Portuguese and so on.

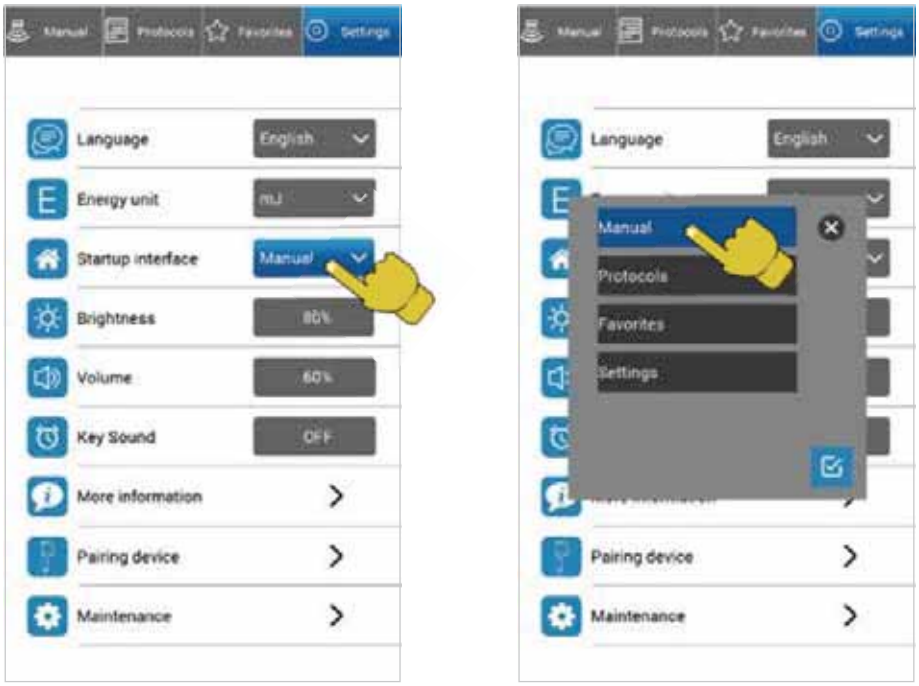
Select your desired language and click the  to confirm.

Energy unit:

Click the 'Energy unit Button' and rotate the knob to select bar, mJ or mJ/mm^2.

Startup interface:

Click the 'startup interface button' and select the start up interface.



Brightness:

Click the 'Light Button' and rotate the knob to adjust the backlight brightness of the main device screen. The range is 10% to 100%, with 10% increments.

Volume:

Click the 'Volume Button' and rotate the knob to adjust the volume of the main device notification sounds. The range is 0% to 100%, with 10% increments.

Key Sound:

Click the 'Key Sound Button' to turn on or turn off the key sound.

More Information Interface:

In this interface, users can:

Information of contraindication:

Press the contraindication to get the access to more information.

Selection of the applicator:

Press the applicator, then you can add the applicators you are going to use in your treatment. Once you set it up, you can switch the selected applicator in the treatment interface by clicking ‘Applicator Button’.
If no applicator is added in More information, the 'Applicator Button' will not be displayed in the treatment interface.

Total shock times:

At the bottom of the interface, view the total shock times of the main device, wired handpiece and wireless handpiece.