

Corstim

Transcranial Electrical Stimulator
for tDCS, tACS, tRNS, Pulse

User Manual

REV. IN-CS 1.1



CE

Transcranial Electrical Stimulator for tDCS, tACS, and tRNS.

This device is in compliance with the requirements of Directive MDD 93/42/EEG and is compatible with the requirements of the following standards: PN-EN 60601-1:2011 + PN-EN 60601-1:2011/A1 Medical electrical equipment -- Part 1: General requirements for basic safety and essential performance and PN-EN 60601-1-2: 2015 Medical electrical equipment -- Part 1-2: General requirements for basic safety and essential performance -- Collateral standard: Electromagnetic compatibility - Requirements and tests as well as the basic requirements of MDD 93/42/EEC.

The product is a medical device.

Corstim is manufactured by QVITI S.A.

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Information about compliance can be obtained from the manufacturer.

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Corstim User Manual

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The device and the software which is supplied with the device belongs to the manufacturer. Upon receipt of the equipment, the user acquires only the right to use the software.

By installing, copying, downloading or otherwise using the Corstim software application that accompanies this device, you agree to be bound by the terms of the End User License Agreement (EULA). If you do not accept these license terms, your sole remedy is to return the entire used device (hardware and software) within 14 days for a refund subject to the refund policy of the distributor of the device.

It is also necessary to seek written permission from the manufacturer before making changes for the use of the device for purposes other than those established.

This guide describes the features on the Corstim device and refers to Corstim device version 1.0.0 and newer. Some functions or accessories described herein may not be available in older versions of the device. Before starting the device, please read these instructions carefully. Keep this document for future reference. Follow warnings and instructions marked on and given by the product.

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1 About Corstim

The Corstim is designed for non-invasive brain stimulation (NIBS) and central nervous system (CNS) stimulation with electrical currents ranging from $\pm 100 \mu\text{A}$ to $\pm 4 \text{ mA}$ (in $1 \mu\text{A}$ intervals) by placing electrodes on the scalp. The device provides stimulation in transcranial direct current stimulation, transcranial alternating current stimulation and transcranial random noise stimulation, (tDCS, tACS and tRNS respectively). Alternatively, Analog Input Mode allows the user to use any analogue signal from an external source. In this case, the stimulation current is adjusted proportionally to the external signal and stimulation is delivered using patient-safe transcranial non-invasive brain stimulation. Using transcranial electrical stimulation, cortical activity can be stimulated or inhibited to modulate plastic changes in the brain. The effects achieved depend on the duration of stimulation, the shape of the current waveform, the current density and the frequency of stimulation. The electrical charge and current density delivered by tES stimulators is well below the charge needed to induce neuron activation and only affects neuron excitability.




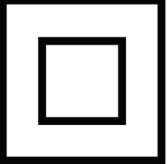




The device is designed for use by professional users (doctors, Rehabilitation, psychologists) or under their supervision. The Corstim stimulator is used in treatment:



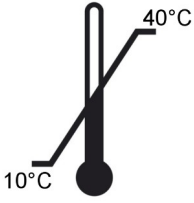
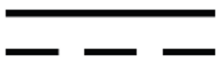


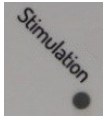
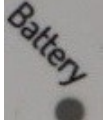

- Neurological disorders:
 - neurological rehabilitation [tDCS],
 - memory disorder [tDCS],
 - aphasia [tDCS],
 - chronic pain [tDCS, tACS, tRNS],
 - migraines [tDCS],
 - Parkinson's disease [tDCS, tACS],
- Mental disorders:
 - depression [tDCS],
 - schizophrenia [tDCS],
 - addictions [tDCS].








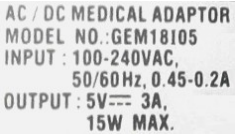
The Corstim, like other stimulators of this type, is not recommended for use in patients with epilepsy or diagnosed anxiety disorders and continuous stimulation should not last longer than 30 minutes.

The Corstim features exceptionally low noise levels, a wide range of parameter settings and a built-in ergonomic user interface for stimulation without the need for external devices such as a computer or tablet.

2 Icons and Labels on Device and Packaging

	<p>Follow the user instructions</p>
	<p>BF applied part. BF applied part is a type B part with additional isolation of the patient circuit from the remaining part of the electrical medical device.</p>
	<p>Special care required. Pay attention.</p>
	<p>Class II device (power unit). Protection of a class II device consists of separating any accessible conductive parts from hazardous parts with double insulation with specified parameters or reinforced insulation to the same level of protection as double insulation. It does not use any earthing measures and is not dependent on the electrical system conditions.</p>
	<p>Manufacturer's address</p>
	<p>Power 'Standby' button. This button does internally not disconnect the device power. Pressing this button turns on or turns off the device.</p>
	<p>Waste electrical and electronic equipment (WEEE) should not be discarded together with other types of waste.</p>
	<p>Serial number</p>

<p>IP41</p>	<p>IP class identification: 4 - protection from solid objects equal or greater than 1.0mm diameter 1 - protected against vertical falling water drop</p>
<p>CE</p>	<p>CE marking constitutes the manufacturer's declaration of the product's conformity with the requirements of Directive 2014/30/WE and 2014/35/UE.</p>
	<p>Handle with care.</p>
	<p>Keep the device dry. Store in a dry place.</p>
	<p>Acceptable storage temperature restrictions (from 10 to 40 °C)</p>
	<p>Direct current (device charging unit)</p>
	<p>Digital in/out triggers terminal (See: 5. Triggers and Analogue I/O)</p>
	<p>USB port (USB port - used only for communication with the device)</p>
	<p>Stimulation - continuously lit (blue) during stimulation and impedance check.</p>
	<p>Battery - continuously lit (yellow) when the device is being charged or green when it is fully charged. The LED goes off when the charger is disconnected.</p>
	<p>Power - continuously lit (green) whenever the device is on.</p>

	<p>Charger terminal 5V DC, max 3A (15W max)</p>
	<p>Medical safety approved (2xMOPP) according to ANSI/AAMI ES60601-1</p>
 <p>EN60601-1 EN60601-1-11</p>	<p>Medical safety approved (2xMOPP) according to IEC60601-1</p>
	<p>For indoor use only</p>
	<p>Polarity of DC power connector</p>
	<p>WEEE; waste electrical and electronic equipment</p>
	<p>FCC Declaration of Conformity</p>
	<p>Mains power unit parameters:</p> <ul style="list-style-type: none"> - AC input power, DC output power, - manufacturer model: GEM18105 medical adaptor, - input power: 100-240VAC at 50/60Hz, - output power: 5V DC, max 3A (15W max)

3 General Device Information

The Corstim is designed for use by or under supervision of specialists.

3.1 Safety, Risks and Prevention of Side-effects

The device is fitted with numerous safeguards to limit the possibility of subjecting participants to any undesired effects. The key variables which may lead to unwanted side-effects include: current dose and the size and positions of electrodes.



Recommended limits of stimulation in tDCS are 2mA of current applied via electrodes with an operating surface of 35cm² in a single session of no more than 30 minutes. The user is required to enter data, prior to stimulation, about the surface area and shape of electrodes being used.

The device measures the current intensity continuously ensuring performance and safety of each stimulation. The device also has a current density control mechanism which is used to adjust the stimulation value and notifies the user of any safe operating limits being reached or exceeded.

The Corstim must not be used on subjects with implanted devices such as a pacemaker and/or deep brain stimulator, as well as intracranial metals such as clippings, coilings, ventriculo-peritoneal shunts and endoprothesis (amongst others) as the device can interfere or cause damage to such metals/devices and/or cause injury to the subject.

An electrostatic discharge may be caused by touching the subject or the device. Such an electrostatic discharge whilst electrodes are attached to a subject's scalp could lead to a mild shock sensation similar to that experienced in everyday life. These currents are not dangerous but can cause an unpleasant sensation. Care should be taken to avoid touching a subject during stimulation.

The most common side-effect occurring as a result of tES is skin irritation, which presents itself in the form of transient redness or, less frequently, as erythema and eschar. To reduce the risk of these side-effects, the use of well moistened salinised sponges under the electrodes or a thick layer of conductive gel is recommended.

Frequently reported experiences for most participants of tES include tingling or a slight burning/itching sensation on the scalp which is caused by skin resistance. This sensation weakens or disappears after approximately 30 seconds. Gradually increasing the current intensity to the target level (fade-in) and a gradual reduction at the end of a stimulation period (fade-out) can help to alleviate these sensations. The Corstim implements a fade-in and fade-out model for power intensity adjustment.

Certain non-specific side-effects have also been described by subjects, such as localised pain and headaches. However, these are not thought to be directly related to stimulation and only appear to occur in subjects who are also suffering from sleep deprivation or hunger.



The risk of inducing an epileptic seizure remains negligible due to the fact that no harmful neuronal hyperpolarization occurs through subcortical stimulation. For safety reasons, subjects who suffer from epileptic episodes are typically excluded from transcranial stimulation studies.

The operator should not touch the subject during stimulation.

The device is designed for transcranial stimulation only. It should not be used for any other types of stimulation, e.g. subdural stimulation. Use of the device for any other purposes may cause tissue damages or may lead to low efficacy of intervention.

Apart from stimulation functionality, device provides analogue input, analogue output, trigger input and trigger output functionalities for interfacing with external devices. Operation of these functionalities is safe for patient and follows requirements of PN 60601-1.

Stimulation of selected narrow areas of the brain could cause short-term interruption of activity in this area, with disruption

of cognitive functions and motor/sensory processes.

The device is fitted with an integrated, rechargeable battery which cannot be removed or replaced. Any attempts to tamper with or remove/replace it can lead to fire, leakage of aggressive substances or explosion. For power unit servicing please contact your distributor.

Any serious incident related to the device must be reported to the manufacturer and the competent authority of the Member State where the user or patient resides



WARNING: Unauthorised opening of the device will void any warranty and may cause electric shock.

3.2 Essential performance, frequently used functions and contraindications

The essential performance of the device is to transcranially stimulate the brain of a subject (patient, research study participant) with given stimulation current parameters for a given amount of time, as defined by the stimulation protocol.

The device may be used for clinical research, comprising of tDCS, tACS, tRNS stimulation or other types of electrical current stimulation through use of the Analog Input Mode.

General contraindications of Corstim use include skin inflammations and epilepsy seizures history. The use of the device in clinical research and the study protocol shall be approved by appropriate entities.

3.3 Device Specifications (Technical description)

Number of channels	1
Stimulation modes	tDCS, tACS, tRNS, Analogue Input Mode
Sham mode	Single-blind and Double-blind
tDCS Parameters	
Stimulation current	max ± 5 mA (increment $1\mu\text{A}$)
Stimulation duration	1-1800s (increment 1s)
Fade-in/out	0-60s (increment 1s)
tACS Parameters	
Amplitude of stimulation current (peak-to-peak value)	max 5 mA (increment $1\mu\text{A}$)
Offset	max ± 3 mA (increment $1\mu\text{A}$)
Stimulation duration	1-1800s (increment 1s)
Fade-in/out	0-60s (increment 1s)
Frequency (carrier)	0.01-600Hz (increment 0.01Hz)
Frequency (modulation)	0-300Hz (increment 0.01Hz)
Modulation index	0-100%
Samples per period	256 (for carrier frequency $< 300\text{Hz}$ and 128 for carrier frequency $\geq 300\text{Hz}$)
Pulse Parameters	
Stimulation current	max ± 5 mA (increment $1\mu\text{A}$)
Stimulation duration	1-1800s (increment 1s)
Fade-in/out	0-60s (increment 1s)
Frequency	0.01-600Hz (increment 0.01Hz)

Signal duty cycle	1-99% (increment 1%, for freq. <50Hz) 10-90% (increment 1%, for freq. ≥50Hz)
tRNS Parameters	
Sample value range	±(200 - 5000)µA (increment 1µA)
Stimulation duration	1-1800s (increment 1s)
Fade-in/out	0-60s (increment 1s)
Noise distribution	rectangular, gaussian
Predefined filters	LP: 100Hz, 250Hz, 640Hz, OFF HP: 50Hz, 100Hz, 250Hz, OFF
Analogue Input Mode Parameters	
Transfer ratio	2mA per V
Current range	±4 mA
Analogue Bandwidth	1kHz (-3dB)
Triggers IN and OUT parameters	
Signal parameters	TTL or 5V CMOS compatible
Other Features	
User programs	1. Creating predefined stimulation programs (PRESET) 2. Directly controlling the protocol through the application interface (API)
Accuracy of current setting	<5% for current <1mA, <1% for current >1mA
Voltage max.	35V
Device rated power	3W
Interface	Touch screen display, control by PC-USB
Digital inputs and outputs	2 digital inputs 3 digital outputs
Analogue Outputs	Main: 0.5V per mA (single ended output) Low level: 2.5mV per mA (differential +, - & GND)
Power supply	Built-in rechargeable battery, total charge typ. 10Ah operating time up to 10h
Mains power adapter (Charger)	Included in the set (Mean Well part number GEM18I05-P1J), 5V/3A DC output 2x MOPP medical safety 80-264VAC 47-63Hz mains input, AC current 0.45A at 115VAC, 0.25A at 230VAC Touch current < 100µA/264VAC
USB	USB micro B plug type, 5V DC, USB 2.0 compatible (12Mbps) (communication only)
Dimensions	150 x 115 x 32mm
Safety control	1. Current density control for a defined electrode; 2. Measurement of impedance (modes: continuous monitoring without stimulation, continuous monitoring with stimulation, single impedance check before stimulation)
Operation of impedance control	Possibility to define a threshold for interrupting stimulation: 10kΩ - 50kΩ Possibility to define user alert threshold: 5kΩ - 15kΩ/off
Electrodes sockets	touchproof DIN42802 1.5mm
Measurement accuracy (stimulation current and voltage)	5%

3.4 Default Parameters Values (Technical description)

Type of setting	Parameter name	Starting value of the parameter
General	Electrode shape	Rectangular
	Maximum impedance	10k Ω
	Impedance alarm	On; 8k Ω
	Sham	None
Electrode rectangular	Dimension a	5cm
	Dimension b	5cm
Electrode round	Diameter	2cm
Electrode shape undefined	Surface area	25cm ²
tDCS	Amplitude	2mA
	Duration	20min
	Fade In	10s
	Fade Out	10s
tACS	Peak to Peak	2mA
	Frequency	10Hz
	Offset	0 μ A
	Duration	20min
	Fade In	10s
	Fade Out	10s
tRNS	Amplitude	2mA
	Highpass filter	None
	Lowpass filter	250Hz
	Noise type	Gaussian
	Duration	20min
	Fade In	10s
	Fade Out	10s
Pulse	Amplitude	1mA
	Frequency	10Hz
	Duty cycle	50%
	Duration	1min
	Fade In	0s
	Fade Out	0s

3.5 Operating, Handling, Transporting and Storage Conditions (Technical description)

The following conditions must be adhered to for the operation, handling and storage of the Corstim. Following these instructions will help to ensure that the device remains functional and minimise any risk of damage to the device or injury to users and subjects.

Keep the device out of and away from:

- Areas where there is a high level of dust, moisture and/or humidity.
- Sources of heat and direct sunlight.
- Strong magnetic or electrical field sources (at least 30cm away).
- Liquid, corrosive chemicals and flammable gasses or liquids.
- Highly pressurised atmospheres (above 1060hPA)

The ambient temperature should be between 10°C – 40°C (50-104°F).

Relatively air humidity should be between 20-95%

Air pressure should be between 700hPA – 1060hPA

Condensation formed in the atmosphere whilst the stimulator is stored, transported or used at low or highly fluctuating temperatures can cause damage to the device. For safety reasons, allow the stimulator to be gradually brought up to room temperature before use (wait minimum one hour before using).

Take care when operating, handling and storing the device to avoid damage to the housing, screen and connection terminals; and to prevent wires being detached.

Do not use the device in a manner other than that outlined in this instruction manual or as advised by the manufacturer. Doing so may damage the device and increase the risk of injury and/or electric shock.

The device is fitted with an integrated, rechargeable battery and is supplied with a mains power unit for charging. Use of a 3rd party power unit other than the one supplied with the device or provided by the manufacturer can cause damage and/or malfunction and increase the risk of injury.

The integrated, rechargeable battery does not need periodic testing or calibration. If the device is stored for more than 12 months, it is recommended to perform re-charging of internal battery.

Do not store the device with less than 50-60% charge in the integrated, rechargeable battery.

Store the device in a secure location, in its original packaging in a cool, dry place free from dust and humidity; and away from unauthorised personnel.

Transport the device in its original packaging free from dust, moisture and humidity. Transport the device in its original packaging, free from dust, moisture, and humidity. Land, sea, and air transport is permissible (battery compliant with UN38.3). Maximum altitude of transport is 2000m.

The device is designed for self-checking its critical functions and will alert if manufacturer's service is required.

There's no need for scheduled maintenance or re-calibration. Any service actions within the device should be performed by the manufacturer's service and engineering department or the authorized and trained representative of the manufacturer only. There are no user-serviceable parts in the device.

The intended life of the product is five years from the date of manufacture.

Dispose the device as required by the legislation of disposal of waste electrical and electronic equipment (WEEE) within the country of use.



WARNING: No modification of this device/equipment is allowed.

3.6 Electromagnetic environment

The intended environment for device's usage is professional healthcare facility environment.

The system the device is certified for consists of the stimulator with two electrodes.

Any cables connected to the triggers (triggers in, triggers out, analog in, analog out) and electrode cables should not be longer than 3 meters.

All interfaces designated as 'outputs' shall be coupled with external high-impedance input ports.

All interfaces designated as 'inputs' shall be driven from low impedance sources.

3.7 Operation

The device is intended for a continuous operation. Practical operation time is limited by batteries capacity.

3.8 Cleaning

The device does not require any sterilisation.

Clean the device with a warm, damp cloth moistened with water and then dry with a separate cloth. Do not use products which contain acetone.

Rubber electrodes with outer sponge should have the cloths removed. The sponges should be washed in 60 degrees water with standard textile washing detergents and rinsed.

Rubbers should be cleaned with isopropanol or disinfectants for medical products (e.g. Schülke Mikrozid AF liquid) afterwards.

When used with rubber electrodes and Ten20 Paste, clean the electrodes with tap water, gently rubbing them. Use isopropanol or disinfectants for medical products (e.g. Schülke Mikrozid AF liquid) afterwards.

Leave the electrodes to dry in room temperature.

If other types of electrodes are in use – refer to user manual of electrodes manufacturer.

3.9 Use



NOTE: The manufacturer or its authorised representative accept no responsibility for any faults and/or injuries a result of misuse of the device.

General recommendations:

- There is no age limitation for subjects undergoing stimulation.
- Settings of the procedure should be consistent with the instructions in this manual.
- Do not expose the device to ionizing radiation, non-ionizing radiation or vibration.
- Use of any accessories or cables other than those supplied with the device or specified in this manual could increase the emission rates or reduce the immunity to external disruption and may consequently cause the device to malfunction.
- Do not put any portable radio communication equipment within 30cm of the device or any of its parts, including the cabling supplied with the device. Otherwise, the device can malfunction.
- The device, when it is in use must not be covered with any materials, such as papers, clothing, notebooks or any type of sheeting.
- If the display is blinking, stop using the device immediately, as you will not be able to control the device functions.
- As a result of significant disruptions caused by such factors as electrostatic discharge, communication with the

device display can be interrupted. When this kind of condition is discovered, the device will refresh the view and the display can blink temporarily. After refreshing, the device will continue working normally. In similar circumstances, battery status indication can be interrupted. In this case, contact your distributor.

- When operating the device, it must be placed on a flat and stable surface, with its display facing upwards, outside the range of the subject's hands, so that the operator can read the messages displayed on screen. The device orientation should facilitate disconnection of applied parts (such as electrodes), and the device should be protected from falling from height onto the operator or subject.
- Do not stack the device on top or under other, as this may cause the device to malfunction.
- If any mechanical damage is found on the device housing, the device should be returned to your distributor for servicing. If any damage occurs during work, interrupt the treatment immediately. Many of these cases involve broken displays.

3.10 Display Messages - Colour Coding

The device displays certain messages concerning its operation to the user.

- Green indicates correct readings.
- Yellow indicates a warning against incorrect stimulation parameters, however without exceeding the preprogrammed safety level.
- Red indicates the highest level of warning, when stimulation cannot proceed or is interrupted if the warning appears during the process. In this case, either the parameters exceed the preset safety level, inadequate settings are used, or the device is inadequately prepared for service (shorted electrodes, too high impedance or impedance $\leq 300\Omega$, charger connected).

Any diagnostic operator with impaired colour perception should pay special attention to the text message and sound signals.

3.11 Package contents

The package contains:

- Main unit (identified by name and serial number)
- Accessories:
 - Mains power adapter with plug adapter (Charger-5V/3A, medical grade Mean Well GEM18I05-P1J)
 - USB cable (MULTICOMP MC002473)
- Optional accessories:
 - Electrode cables (Spes Medica BEC152126S12 or equivalent)
 - Reusable silicon electrodes (Spes Medica SIL0005050 or SIL0005070 or equivalent)
 - Disposable electrode support cap (Spes Medica D00000SA000 or equivalent)



NOTE: Use only the original charger supplied by the manufacturer. Use of a non-authorized charger may result in a device malfunction or unacceptable risk.

3.12 Device Layout

Figure 1 shows the front view of the device, with the following components:

- 6 Charger terminal
- 7 Power 'Standby' button

- 8 Device status LEDs
- 8.1 Power - continuously lit (green) whenever the device is on.
- 8.2 Battery - continuously lit (yellow) when the device is being charged or green when it is fully charged. The LED goes off when the charger is disconnected.
- 8.3 Stimulation - continuously lit (blue) during stimulation and impedance check.
- 9 USB communication port
- 10 Digital in/out triggers terminal

The markings and labels are readable from 30 cm by a person without a visual impairment.



Figure 1. Corstim front view

Figure 2 shows the top view of the device, including its 4.3" touch screen user interface (GUI).

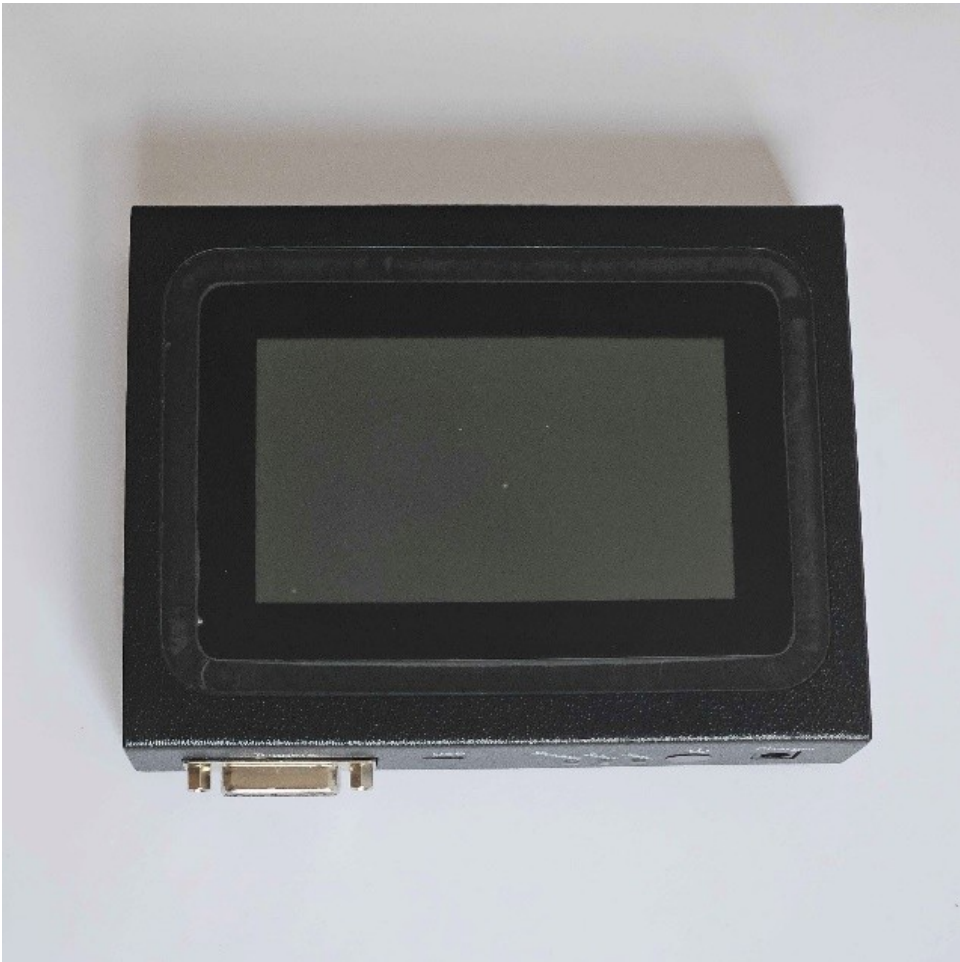


Figure 2. Corstim top view

Figure 3 shows a view of the device from the left, with visible 1.5mm terminals for red electrode - anode (+) and black electrode - cathode (-).



Figure 3. Left side view of the Corstim device.

3.13 Power Supply and Charging

Power supply to the Corstim is provided from an integrated lithium rechargeable battery. The battery may not be fully charged on delivery and should be charged before first use. The charging status is shown as a percentage in the upper part of the screen.

When the battery charge is low (<5%) the device displays a low battery screen containing a relevant message. The device will automatically turn itself off after 10 seconds (Figure 4).

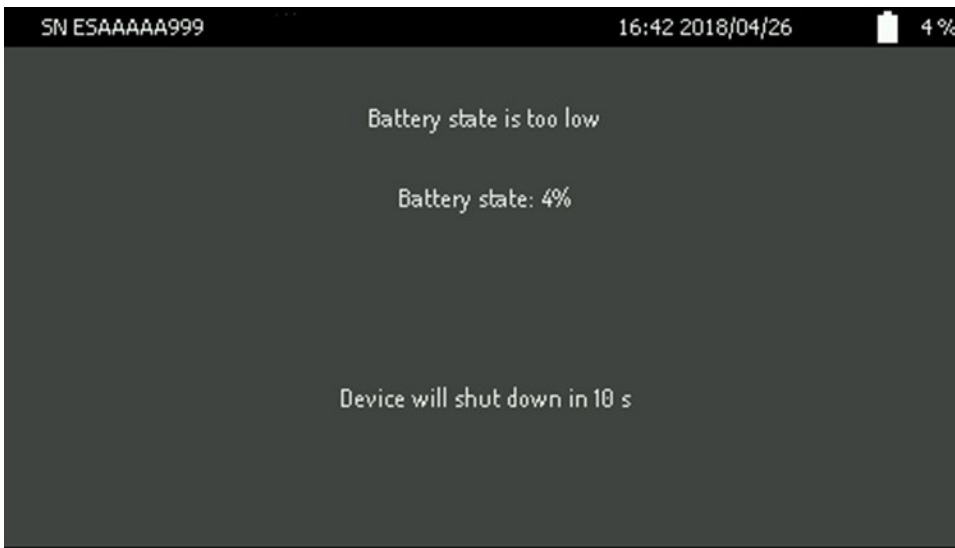


Figure 4. Battery state is too low.



Figure 5. Proper connection of power supply

To charge the battery, connect the power unit supplied with the device to the charger terminal (Figure 5).

A power unit connected to the device will be detected automatically and a charging screen will be displayed (Figure 6). After 10s the device is shut down automatically. Charging is indicated by the Battery LED which will be continuously lit. A yellow LED means the device is being charged and green indicates when it is fully charged. The LED goes off when the charger is disconnected. The charging screen displays whether the charger is connected/disconnected, the charging percentage, the charging process, and time left to device shutdown (seconds).



As a safety precaution, the device cannot be charged and operated at the same time. The user or subject must not touch the electrodes while the charger is connected or while the device is charged.

Allow approximately 10 hours to fully charge the battery.

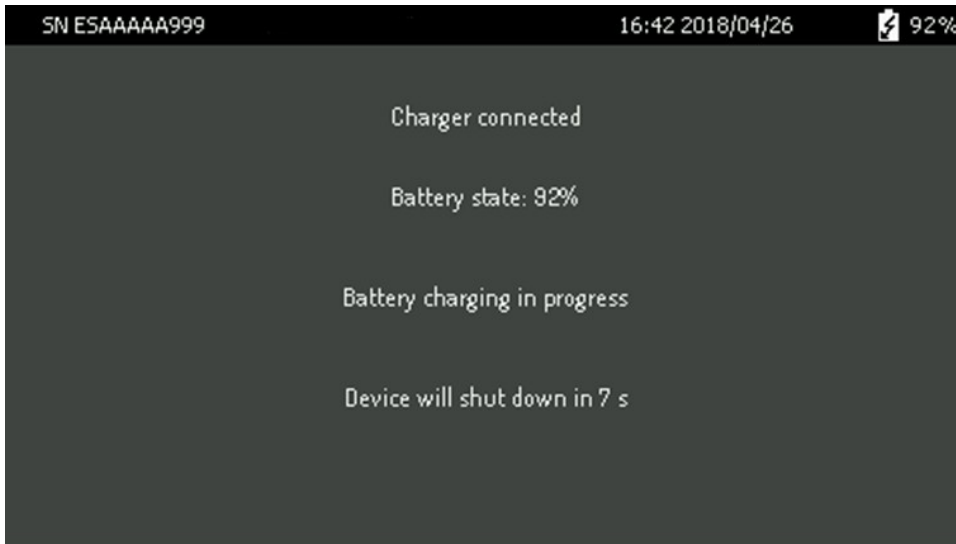


Figure 6. Charging screen

3.14 Starting the Device

To start the device, press the Power button (Figure 1) and hold for approx. 2 seconds, or until the green Power LED goes on (Figure 1). A welcome screen will be shown on the device.

3.15 Switching Off the Device

To switch off the device, press and hold the Power button for approx. 2 seconds or until the green Power LED goes off. You can shut down the device after making sure that the stimulation process is over.

3.16 Storing Last Settings

The last custom settings of the user will be stored at shutdown and restored at the next start-up.

3.17 Connecting Electrodes

Electrodes should be connected to their corresponding electrode terminals (Figure 7) using 1.5mm pin cables. To disconnect an electrode, hold the pin and pull.



Never remove electrodes while stimulation is in process. For safety reasons, it is recommended that the cables be plugged into the correct inputs according to their colour - insert the red cable into the red socket (Figure 7).



Figure 7. Correct electrodes cables connection



NOTE: All electrodes and cables must have a CE declaration of conformity to be approved for use with this device.

Rubber carbon electrodes or sponge electrodes are acceptable if they are specifically manufactured for use with transcranial electric stimulator devices. Spes medica 50x70 mm electrodes are recommended (Figure 8). Before use, check that the electrodes are dedicated to transcranial electrical stimulation and are not broken.

Special conductive gel should be applied to the electrodes (Figure 9). Never use tap water to wet electrode sponges.

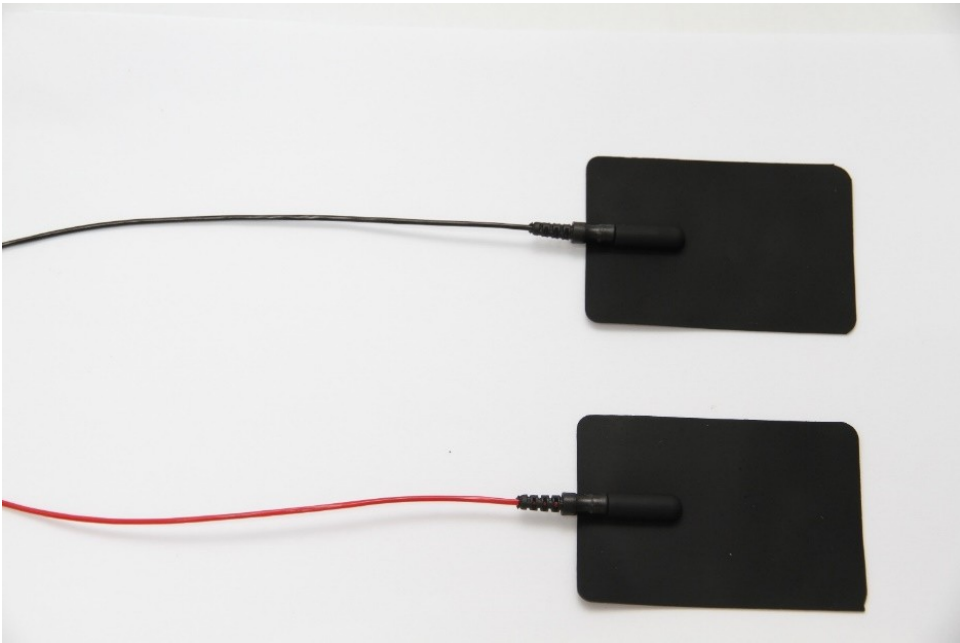


Figure 8. Electrodes connected



Figure 9. Arrangement of the gel on the electrodes.

3.18 Impedance Control

The unit is fitted with an integrated impedance control mechanism. Impedance measurement begins before the start of every stimulation and continues during the stimulation. In addition, users can initiate impedance measurement from the stimulation summary screen. An acceptable impedance threshold and impedance alert level can be programmed into the device (see: 4.11.2 and 4.11.3). Stimulation is not possible when the impedance value exceeds the threshold at 50k Ω .

In addition, stimulation will stop when impedance value falls below 300 Ω , indicating a shorting in the electrodes. The impedance control function will also verify whether the preset values can be reached with the current level of impedance. To control impedance, the device generates a low-intensity electric signal. For details, contact your distributor.

4 Graphic User interface (GUI), Parameter Selection and Stimulation

The Corstim device has been designed with a unique graphic user interface which means that it can be controlled directly via the integrated touch screen. This makes it ideal for studies that require mobile tES.

4.1 Welcome Screen

On start-up the device will display the Welcome screen.

The upper bar of the screen contains the device serial number, operating mode, date and time, and battery status. The Corstim logo is displayed in the centre.

After a few seconds, the device will automatically switch to the Home screen.

4.2 Home Screen

On the Home screen (Figure 10) four buttons are available for the user.

- Custom stimulation – allows selection of the stimulation method (parameters);
- Presets – where predefined stimulation settings can be selected.
- Analogue Input Mode – allows special stimulation mode controlled via external analogue source
- Settings – where the device date and time can be change and where stimulation logs can be managed.

To setup a new stimulation select 'Custom Stimulation'.

To start a previously saved preset select 'Presets'.



Figure 10. Home screen

4.3 Settings

Press 'Settings' to redirect to the Settings screen (Figure 11).

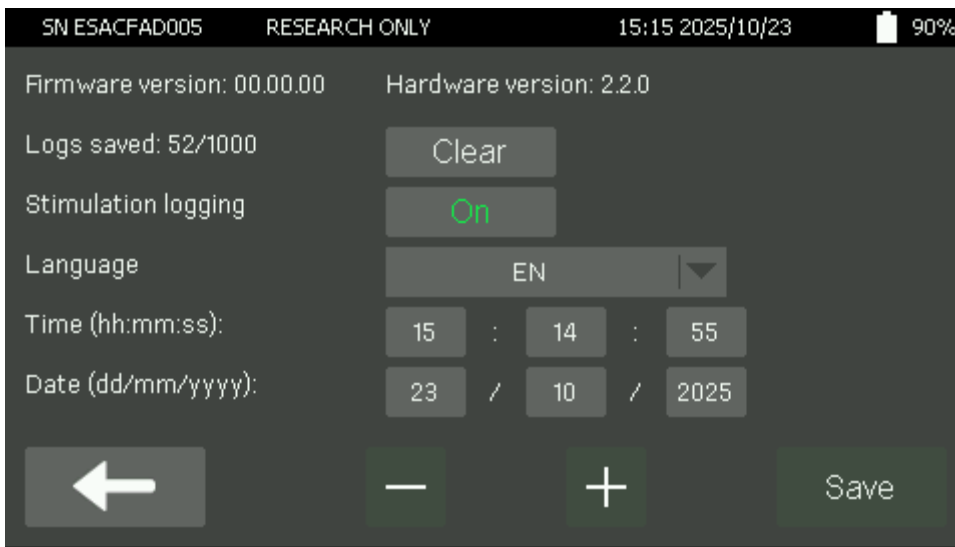


Figure 11. Settings screen

The Settings screen displays information about the software and hardware versions. Below that is information about the number of session logs stored in the device's internal memory. Each log contains information about every started stimulation (stimulation parameters, time, settings, measurement, and stimulation end status). Logs are recorded even when the device is used without a computer and application. The device can store up to 1000 logs. Logs can be downloaded and deleted from the device using a dedicated application. Logs can also be deleted using the "Clear" button. Logging can be disabled using the "On/Off" button.

The device allows changing the user interface language. English and Polish are available. The interface language setting does not affect communication with a PC or the language set in the application.

From the settings screen, it is possible to set the device's internal clock (RTC). To change the time and date settings, press the appropriate button and set the desired value using the + and - buttons. Time and date settings are confirmed by pressing the "Save Date" button. The clock starts counting time immediately after saving the settings.

The format of the time and date setting is set out below. To edit these on the device select the parameter you want to change.

- hh – hours,
- mm – minutes,
- ss – seconds,
- dd – day,
- mm – month,
- yyyy – year.

Pressing + or – will either increase or decrease the value. Press 'Save Date' to save the new time and date. Pressing back arrow will move to home screen but without saving date and time.

4.4 Custom Stimulation Screen

When 'Custom Stimulation' is selected, the stimulation setup screen is displayed (Figure 12).

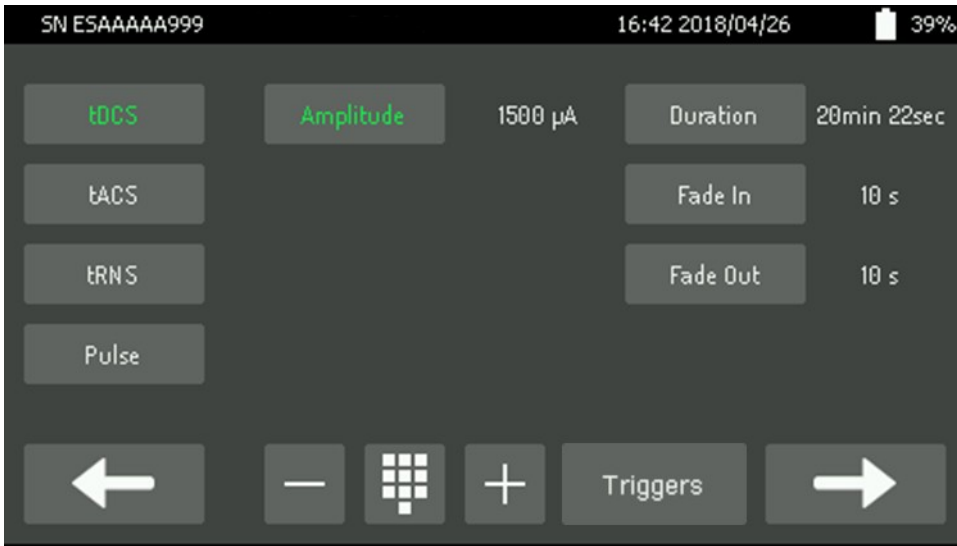


Figure 12. Custom Stimulation screen (tDCS)

Available stimulation modes are listed as buttons in left column. The currently selected stimulation mode is highlighted in green. To switch the stimulation mode, select the desired mode by pressing the relevant button.

To return to previous screen, press the left arrow on the bottom of the screen.

From the stimulation parameters settings, you can access the external control signal settings, which are described in detail in a separate section of the manual (5 Triggers and Analogue I/O).

When changing the stimulation mode, different sets of parameters are available, as shown below. (Figure 12, Figure 13 and Figure 14)



Figure 13. tACS highlighted showing different parameters.

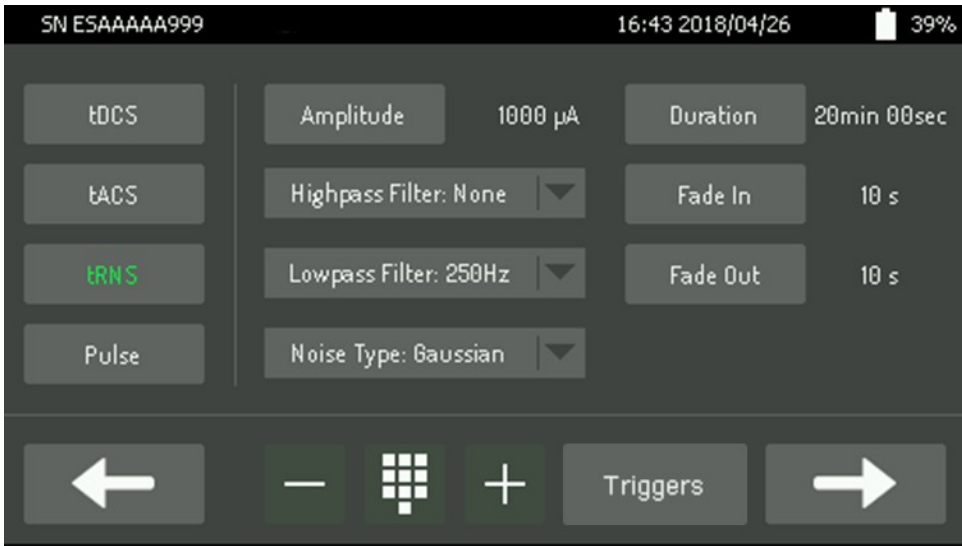


Figure 14. tRNS highlighted showing different parameters.



WARNING! On some devices, certain stimulation modes (modalities) may be inactive (Figure 15). This also applies to the Analog Input mode on the start screen (Figure 16). To unlock unavailable stimulation modes, please contact the distributor.

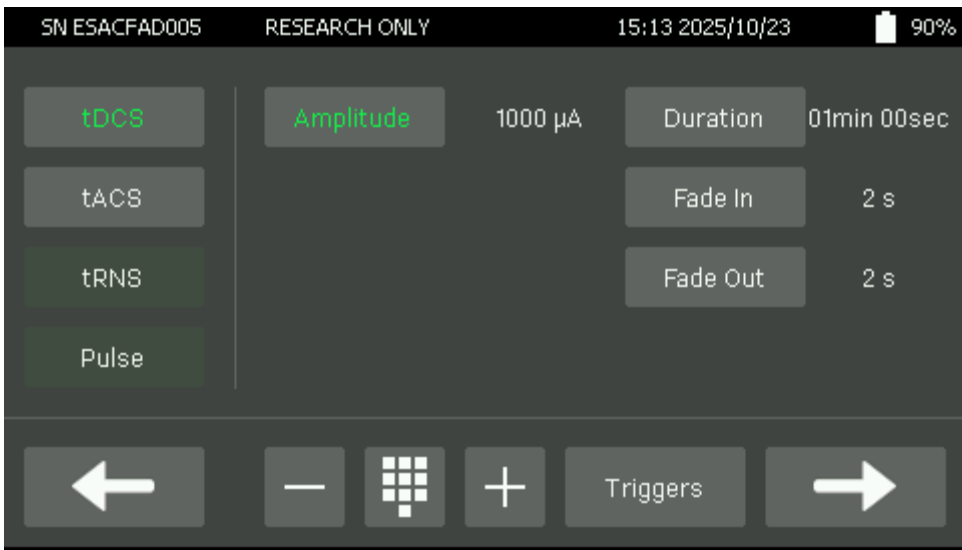


Figure 15. tRNS and Pulse locked.




Figure 16. Analog Input Mode locked.

4.5 Editing Stimulation Parameters

To customize the stimulation, select the desired stimulation method. The method will be highlighted in green once selected.

There are three ways of editing parameters:

- + and – to increase/decrease parameter value
- Keypad – when the keypad button  is pressed – the desired levels can be entered via the keypad (Figure 17).
- Dropdown menu – some parameters have a fixed number of options selected from a dropdown menu.

To start editing parameter values, first press button/dropdown item with its name. Then use one of the methods to set new value.

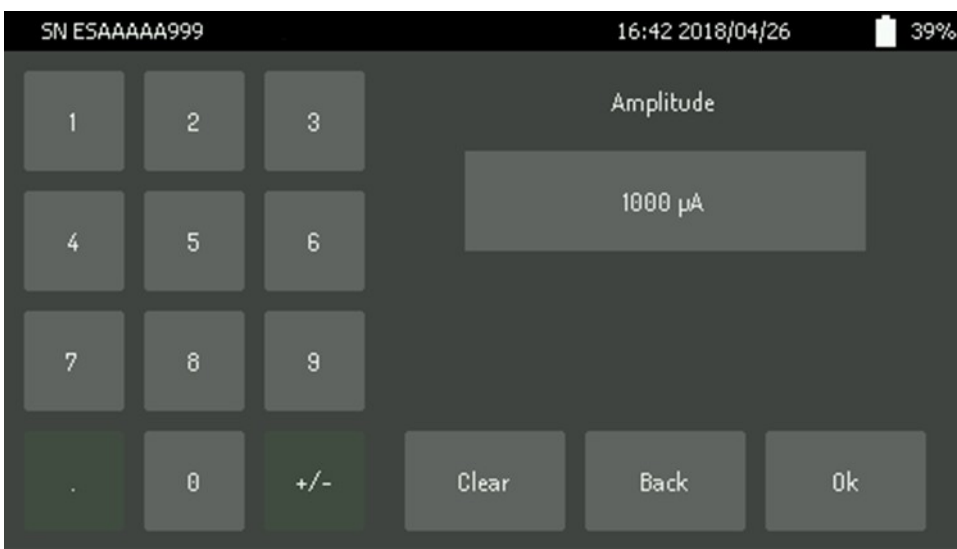


Figure 17. Amplitude entered using the keypad.

4.6 General Settings for all Methods of Stimulation

Setting the stimulation time (excluding duration, fade-in and fade-out) for all methods of stimulation is identical (Figure 18). See part 4.6.1 and 4.6.2.

Other parameters differ depending on the selected type of stimulation (see part 4.7, 4.8 and 4.9.).



Figure 18. tACS mode settings.

4.6.1 Duration

Duration is a parameter that determines the time of a proper stimulation with a given current value (Figure 19). The units of this parameter are minutes [min] and seconds [sec]. The highest possible value to be set for this parameter is 30mins and the lowest possible value is 1sec.

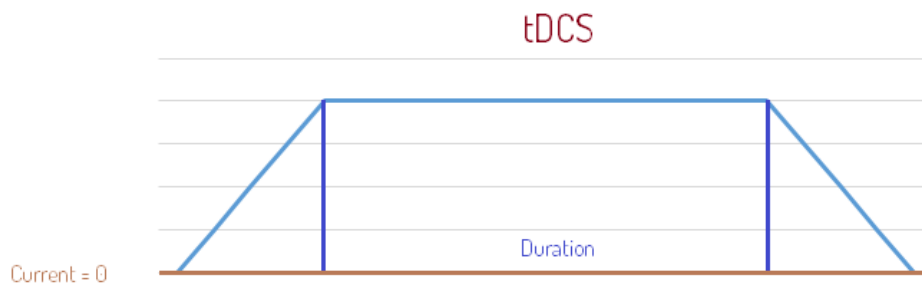


Figure 19. Duration.

To edit, select the stimulation mode and then press “Duration.” Next, press the keyboard button and enter the desired number of minutes (Figure 20). If a value exceeding the allowable maximum is entered, it will automatically be adjusted to the highest permitted value. If the maximum number of digits for minutes is entered, the numeric keyboard will be deactivated, but it will still be possible to enter seconds by pressing the “sec” button. If it is necessary to set the time with second-level accuracy, press the “sec” button on the keyboard and enter the desired number of seconds (Figure 21). To confirm, press OK. If a mistake was made, a new value can be entered after pressing the “Clear” button. To cancel the new setting and keep the previous value, press “Back.” The stimulation time can also be set using the + and - buttons, which increase/decrease the setting by 1 second.

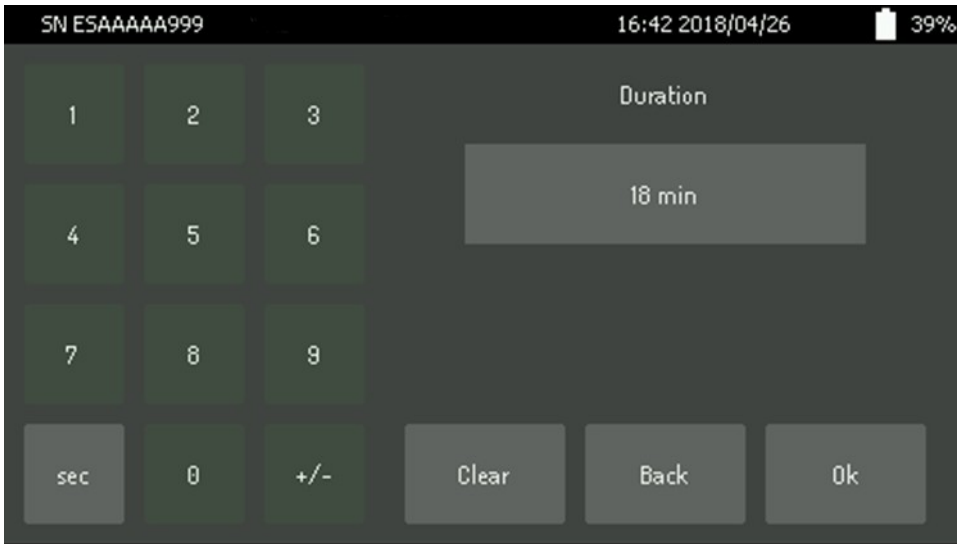


Figure 20. Maximum number of digits for minutes of stimulation reached. Press 'sec' to increase duration by seconds.

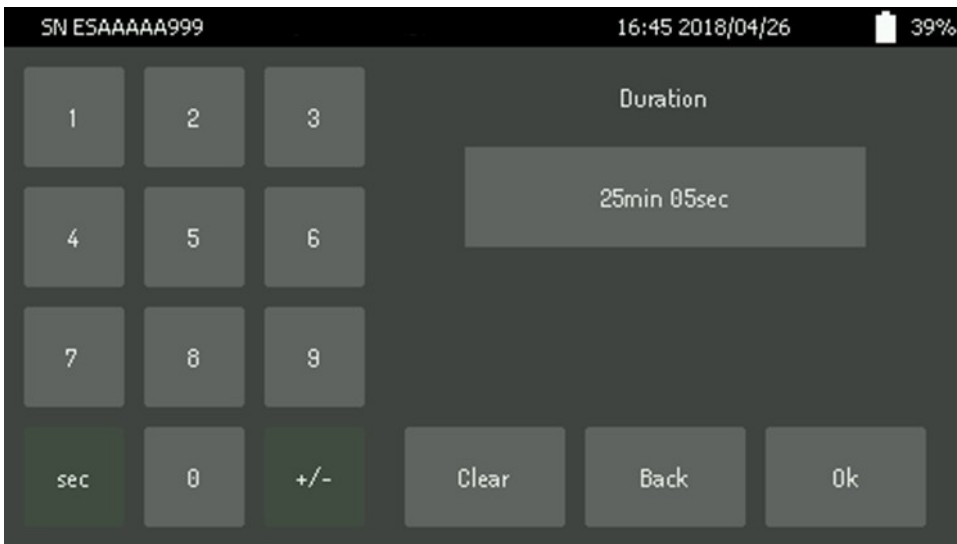


Figure 21. Duration setting screen.

'Ok' is available when an appropriate value is entered.

'Back' closes the keypad and return to the previous screen without saving.

'Clear' erases entered value so that a new value can be entered.

4.6.2 Fade-in and Fade-out

Fade-in and Fade-out is a time value for the signal to rise before stimulation and fall after the stimulation has finished (Figure 22). This is a key safety feature and is a systematic increase of the current so that it will not burn the subject. The unit of this parameter is a second [s]. Fade-in and Fade-out can be set between 0-60 seconds.

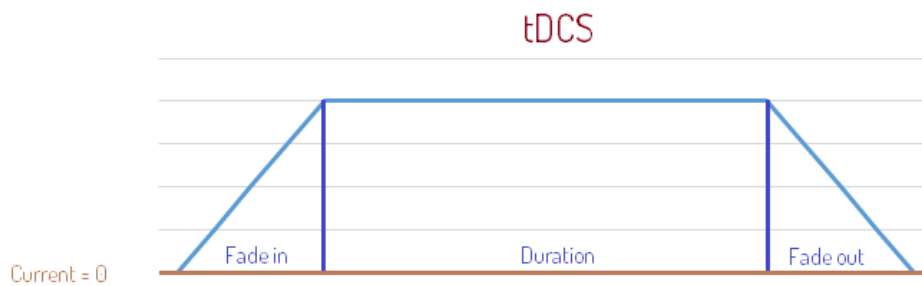


Figure 22. Fade in and fade out

Fade-In or Fade-Out can be edited once the stimulation method had been selected. The + and – buttons will increase/decrease by 1 second the duration of the Fade-In or Fade-Out, which will be highlighted in green to show which is being edited.

Pressing the keypad icon will bring up the keypad screen (Figure 23).



Figure 23. Fade in setting screen.

Should a value be entered that exceeds the permitted maximum, it will be automatically corrected to the highest permissible value. If the maximum number of digits is entered, the numeric keypad is deactivated.

‘Ok’ is available when an appropriate value is entered.

‘Back’ closes the keypad and return to the previous screen without saving.

‘Clear’ erases entered value.

4.7 Specific tDCS Parameters

tDCS (transcranial direct current stimulation) is a form of stimulation that uses direct current.

4.7.1 Amplitude

Amplitude is the only parameter for tDCS. It represents the value of the tDCS electrical current (Figure 24). The unit of this parameter is a microampere [μA]. The value range for Amplitude is between $100\mu\text{A}$ and $4000\mu\text{A}$.

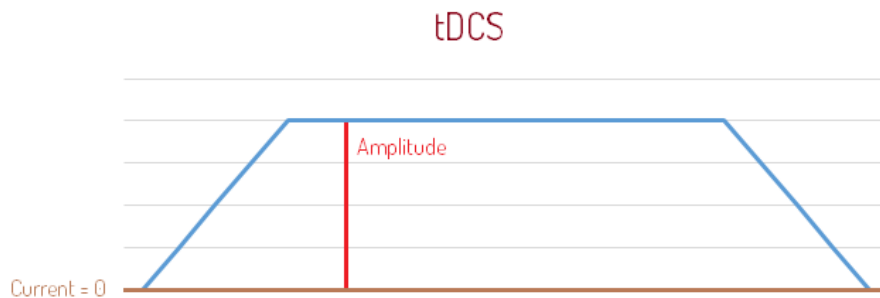


Figure 24. tDCS Amplitude.

To edit the amplitude parameter, press the 'Amplitude' button. Use the + and - buttons to change amplitude levels at increments of 100 μ A. Pressing the keypad will access the keypad screen, where exact amplitude levels can be entered (Figure 25).

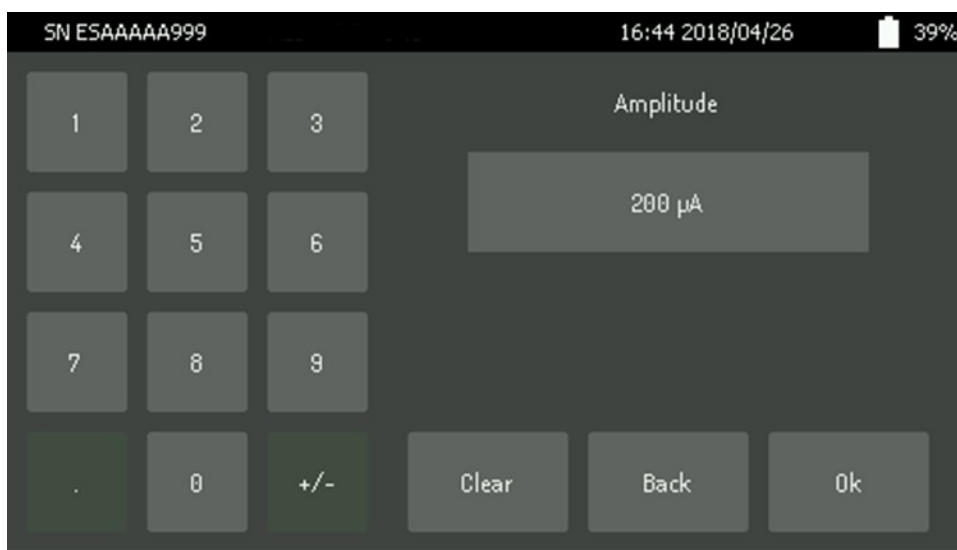


Figure 25. Amplitude setting screen using the keypad.

Should a value be entered that exceeds the permitted maximum, it will be automatically corrected to the highest permissible value. If the maximum number of digits is reached the numerical buttons will be deactivated.

'+/-' changes current polarity (if applicable).

'Ok' is available when an appropriate value is entered.

'Back' closes the keypad and returns to the previous screen without saving.

'Clear' erases the entered value.

4.8 Specific tACS Parameters

tACS (transcranial alternating current stimulation) conducts a sinusoidal current at a chosen frequency, amplitude and offset. Amplitude modulation signal is also available.

4.8.1 Amplitude

Amplitude is the change between peak (highest signal value) and trough (lowest signal value, which can be negative). (Figure 26)

To edit the Amplitude parameter go to section 4.7.1.

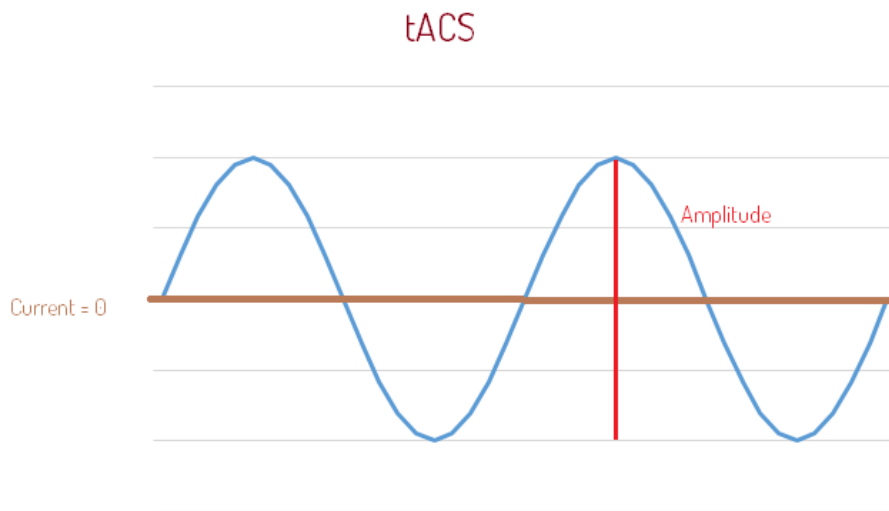


Figure 26. tACS Amplitude.

4.8.2 Frequency

The Frequency parameter in tACS is the number of cycles of the periodic phenomenon occurring in a unit of time (Figure 27).

The unit of this parameter is a hertz [Hz]. The value range for this parameter is between 0.01Hz and 600Hz.

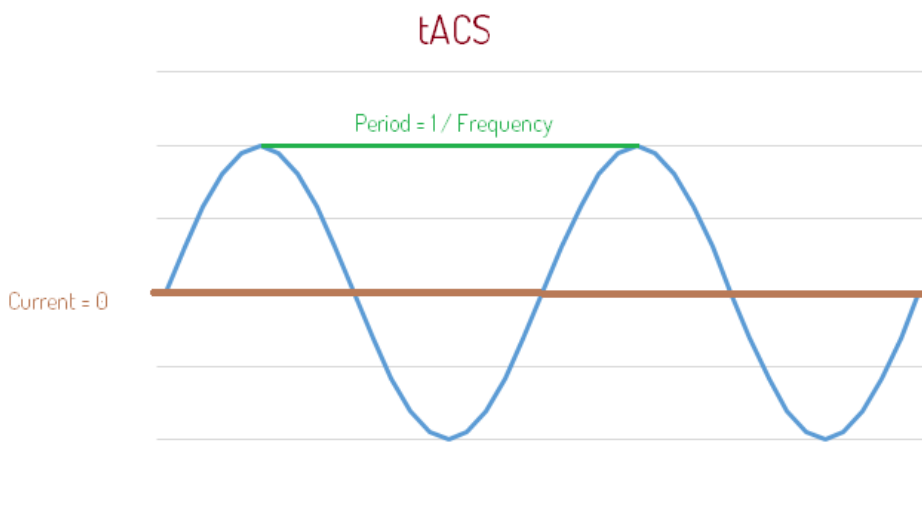


Figure 27. tACS Frequency.

To edit the 'Frequency' parameter in tACS mode, use + and - to increase/decrease the frequency in increments of 10Hz. Pressing the keypad will access the keypad screen, where exact frequency levels can be entered (Figure 28).

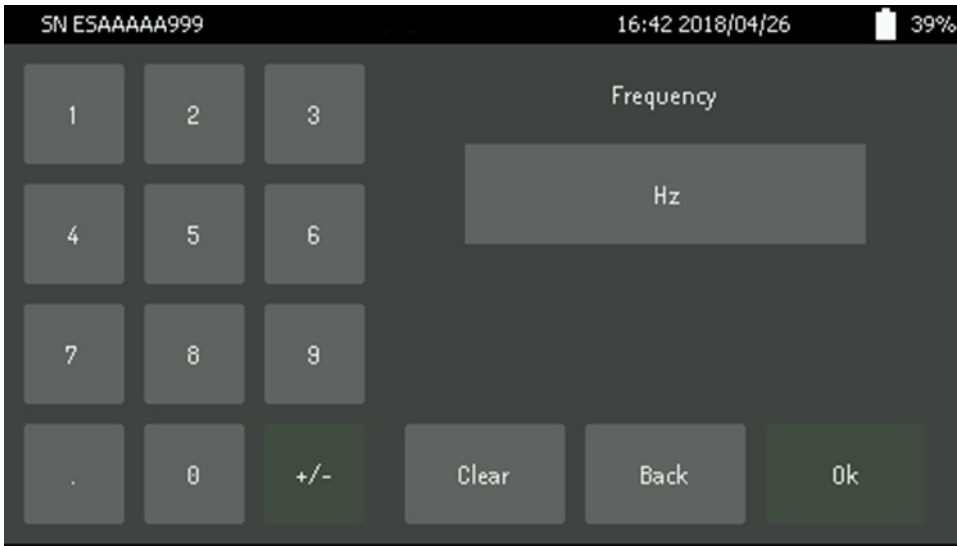


Figure 28. Frequency setting screen.

To enter a fractional value, press the decimal button. Should a value be entered that exceeds the permitted maximum, it will be automatically corrected to the highest permissible value. If the maximum number of non-fractional digits is reached, the numeric keypad is deactivated, but the user can add 2 decimal places after pressing the decimal button (Figure 29).

'Ok' is available when an appropriate value is entered.

'Back' closes keypad and returns to the previous screen without saving.

'Clear' erases the entered value so that a new value may be entered.

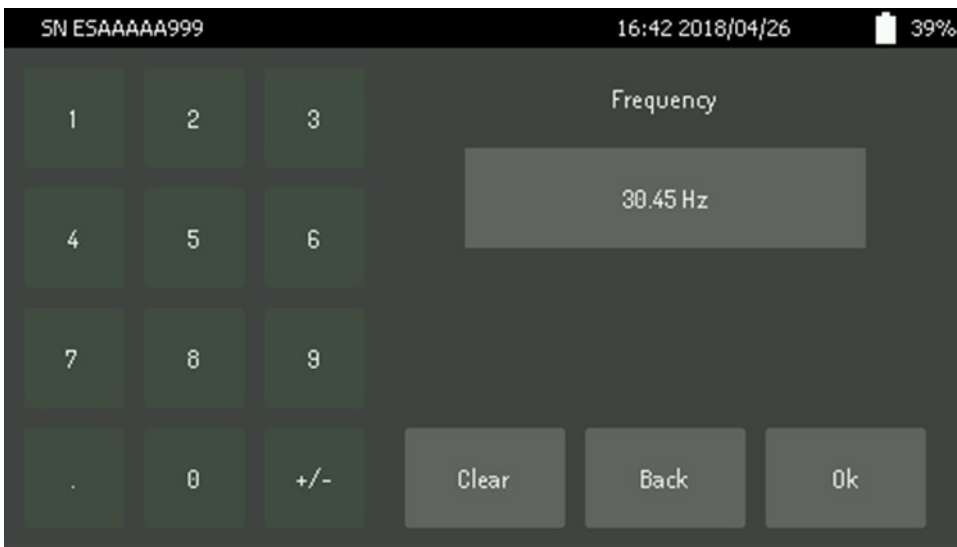


Figure 29. Fractional part of frequency parameter.

4.8.3 Offset

Offset is the movement of the entire sinusoid by a given value along the axis of value. In practice, this means giving a constant mean value to the tACS stimulation (Figure 30).

The unit of this parameter is a microampere [μA]. The value range for this parameter is between minus $3000\mu\text{A}$ and $3000\mu\text{A}$. Should a value be entered that exceeds the permitted maximum, it will be automatically corrected to the highest permissible value.

tACS

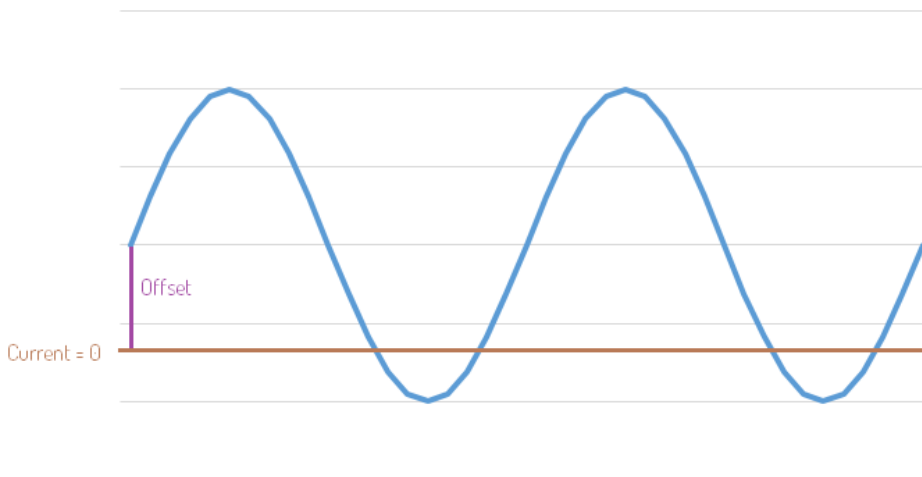


Figure 30. tACS Offset.

The offset parameter is only available for tACS.

To edit Offset parameters, press 'Offset' and use + and – to increase/decrease values by 100 μ A. Pressing keypad will bring up keypad screen (Figure 31).

The sum of Offset and Amplitude values cannot exceed the maximum permitted current (it has to be in the range of $\pm 4000\mu$ A). Should the entered value cause a violation of that restriction, the other parameter is decreased to avoid it.



Figure 31. Offset setting screen.

'+/-' changes offset polarity.

'Ok' is available when an appropriate value is entered.

'Back' closes the keypad and returns to the previous screen without saving.

'Clear' erases entered value so that a new value may be entered.

4.8.4 Envelope Amplitude

Envelope amplitude is the depth of modulation of the tACS signal, which changes the value of the amplitude given to the

stimulation (Figure 32).

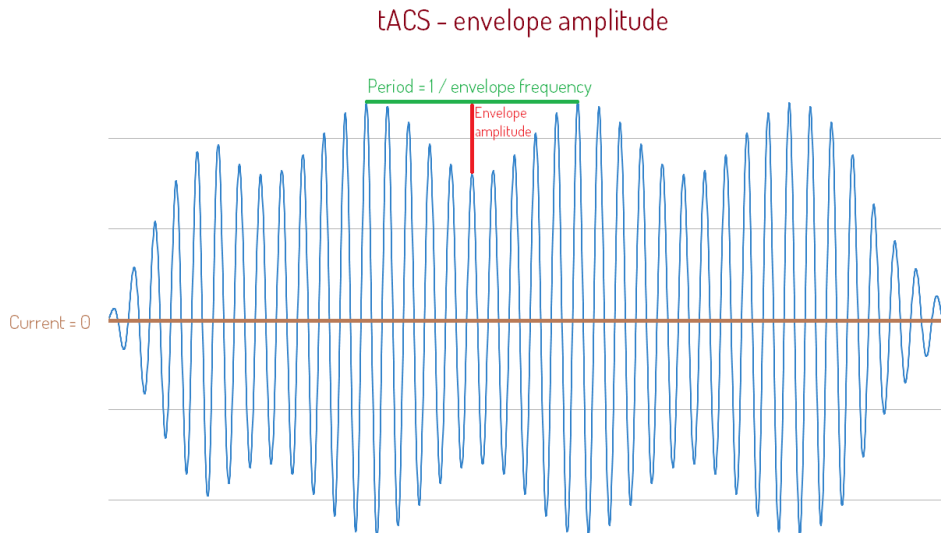


Figure 32. tACS Envelope amplitude and frequency.

The unit of this parameter is a microampere [μA]. The envelope amplitude value shall not be bigger than half of the stimulation amplitude.

To edit, press 'Env. ampl.' in tACS mode. Use + and – to increase/decrease the value by $100\mu\text{A}$. Pressing keypad will bring up keypad screen (Figure 25).

The value of envelope amplitude cannot exceed half of the amplitude of the carrier. Should the entered value cause a violation of that restriction, the envelope amplitude is decreased to avoid it.

'Ok' is available when an appropriate value is entered.

'Back' closes the keypad and returns to the previous screen without saving.

'Clear' erases entered value so that a new value may be entered.

4.8.5 Envelope Frequency

Envelope frequency is the frequency of modulation signal (Figure 32).

The unit of this parameter is a hertz [Hz]. The value range of this parameter is between 0Hz (where no modulation will occur) and 100Hz. The value of envelope frequency cannot exceed the half of the carrier frequency (tACS frequency). Should the entered value cause a violation of that restriction, the envelope frequency is decreased to avoid it.

To edit, press 'Env. freq.' in tACS mode. Use + and – to increase/decrease the envelope frequency by 10Hz. Pressing keypad will bring up keypad screen (Figure 28). The envelope frequency value shall not be bigger than half of the stimulation frequency.

'Ok' is available when an appropriate value is entered.

'Back' closes the keypad and returns to the previous screen without saving.

'Clear' erases entered value so that a new value may be entered.

4.9 Specific tRNS Parameters

tRNS (transcranial random noise stimulation) mode utilises alternate current along with given maximum current value, spectrum and distribution.

4.9.1 Amplitude

The Amplitude parameter in tRNS mode defines the current range from which all samples will be generated (peak-to-peak value). (Figure 33)

Setting any filter or setting Gaussian rather than rectangular noise type may decrease (or under some circumstances even increase) the actual amplitude due to a random nature of the signal.

To edit the amplitude parameter go to section 4.7.1.

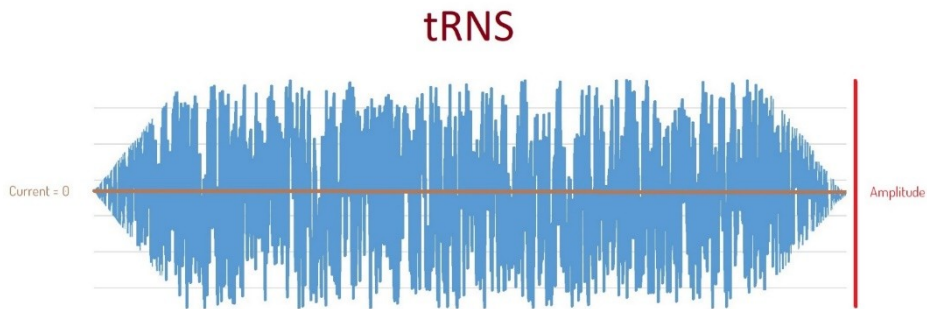


Figure 33. tRNS Amplitude.

4.9.2 Highpass Filter

The Highpass Filter is available specifically for the tRNS signal. A highpass filter attenuates frequencies below the selected value. Edit the Highpass Filter cut off frequency by selecting the required cut-off frequency from the Highpass Filter drop down menu (Figure 34).

Note: A highpass filter cannot have a cut-off frequency equal to or higher than that of the lowpass filter.

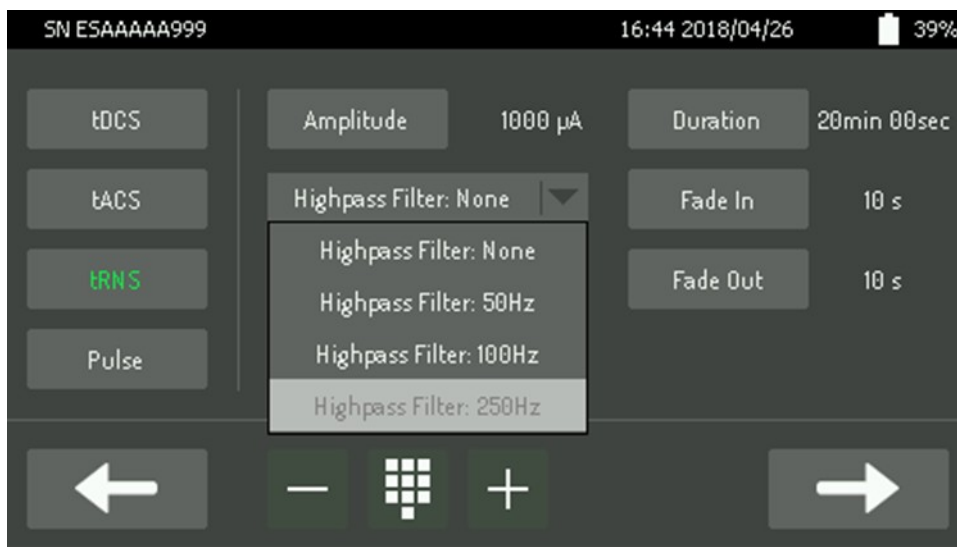


Figure 34. Setting highpass filter for tRNS.

4.9.3 Lowpass Filter

The Lowpass Filter is available specifically for tRNS Signal. A lowpass filter attenuates frequencies above the selected value. Edit the Lowpass Filter cut off frequency by selecting the required cut-off frequency from the Lowpass Filter drop down menu (Figure 35).

Note: The lowpass filter cannot have a cut-off frequency equal to or lower than that of the highpass filter.



Figure 35. Setting lowpass filter for tRNS.

4.9.4 Noise Type

There are two types of noise distributions available for tRNS.

- Rectangular – causes samples to be generated evenly over the whole signal current range.
- Gaussian – weights samples more to the middle of the current range than at either extremity.

To choose a noise type, first select the desired tRNS method, then select the desired noise type from the two options available (Figure 36).

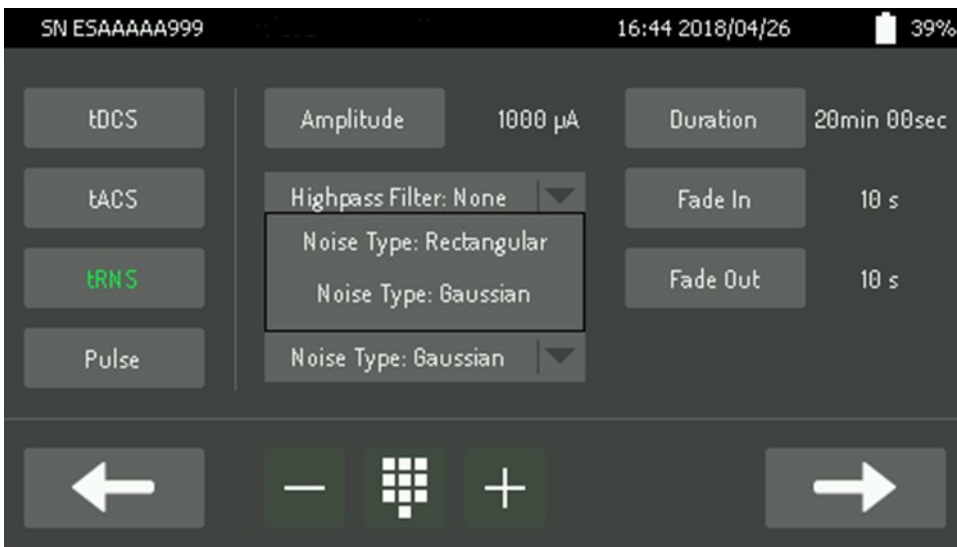


Figure 36. Noise type selection dropdown.

4.10 Specific Pulse Parameters

The Pulse mode uses a current in the form of monopolar rectangular pulses with a specified amplitude, frequency, and duty cycle. These parameters are shown in Figure 37 i 38.

4.10.1 Amplitude

The pulse amplitude in Pulse mode refers to the current value during the active part of the pulse. The unit of this parameter is micro ampere [µA]. The range of this parameter is from 200µA to 5000µA.

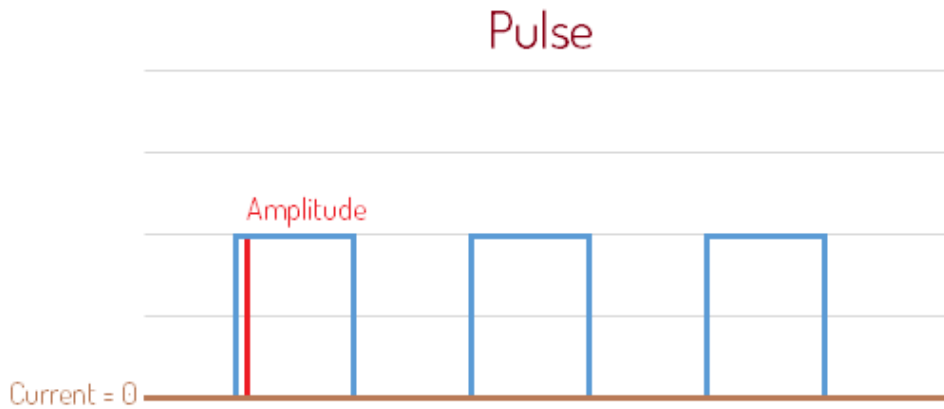


Figure 37. Amplitude in Pulse mode.

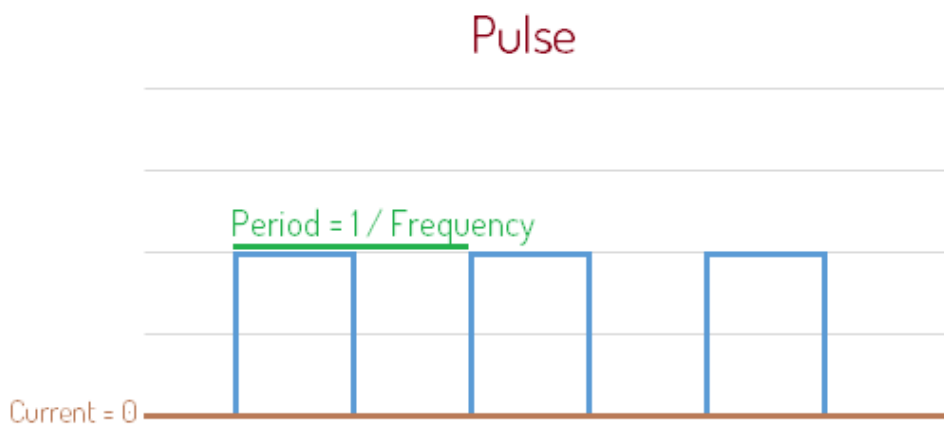


Figure 38. Frequency in Pulse mode.

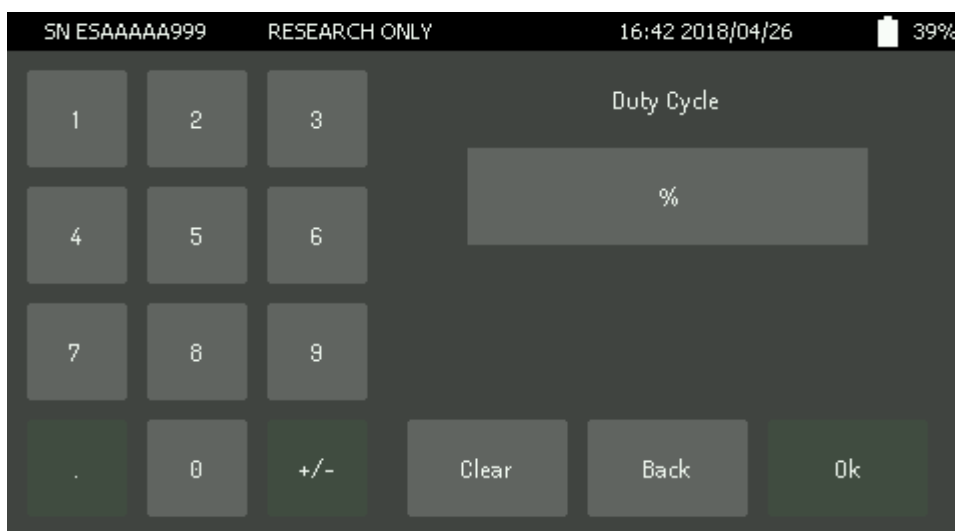


Figure 39. Duty cycle settings screen in Pulse mode.

4.10.2 Frequency

The pulse frequency determines the number of signal pulses per second. The unit of this parameter is hertz [Hz], and the

available range is from 0.01Hz to 600Hz. The frequency parameter can be edited as described in section 4.8.2.

4.10.3 Duty cycle

This parameter describes the percentage of the active signal (non-zero current) within one signal period. The unit of this parameter is percent [%], and the available range is from 1% to 99%. For frequencies equal to or higher than 50Hz, the Duty Cycle is limited to the range of 10% to 90%.

To edit, press “Duty cycle” in Pulse mode. Use the + and – buttons to increase or decrease the value by 5%. The parameter value can also be entered using the keypad. (Figure 39).

4.11 Advanced options

Once all stimulation parameters have been set, press the right arrow button at the bottom right corner of the screen to proceed to the advanced options screen.

The following options are not directly related to the signal to be used in stimulation. However, it is still necessary that they are set.

4.11.1 Electrodes

This option relates to the type of electrode to be used in the next stimulation. The surface area of the electrode is calculated using provided parameters so that a current density calculation can be made. This step can be skipped; however, for safety reasons skipping this step is not recommended.



Note: If your electrodes are of different sizes, always use the size of a smaller electrode.

To choose an electrode type, select the electrode type dropdown menu and choose the desired option (Figure 40). There are 4 types of electrode shape possible:

- rectangular
- round
- shape undefined
- ignore electrode size

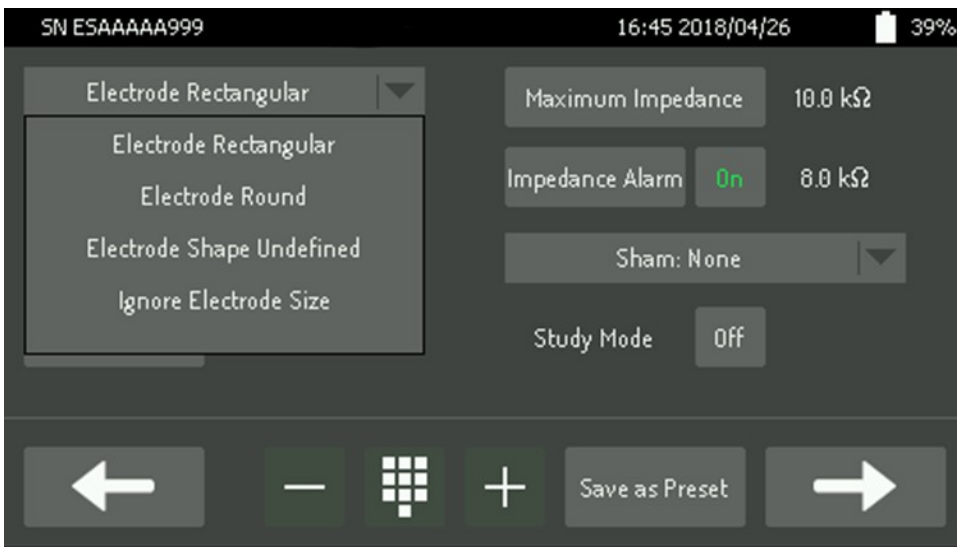


Figure 40. Electrode type dropdown shown at the top left.

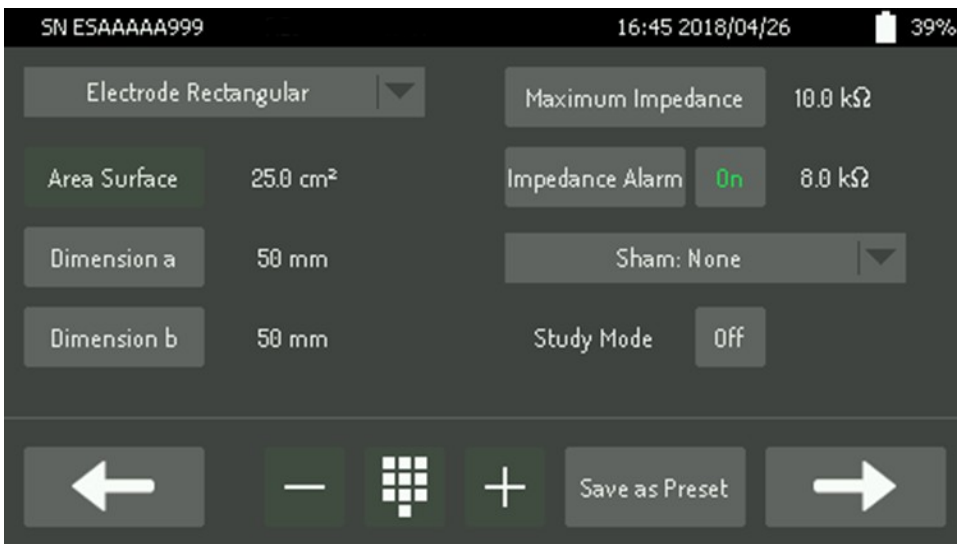


Figure 41. Electrode Rectangular.

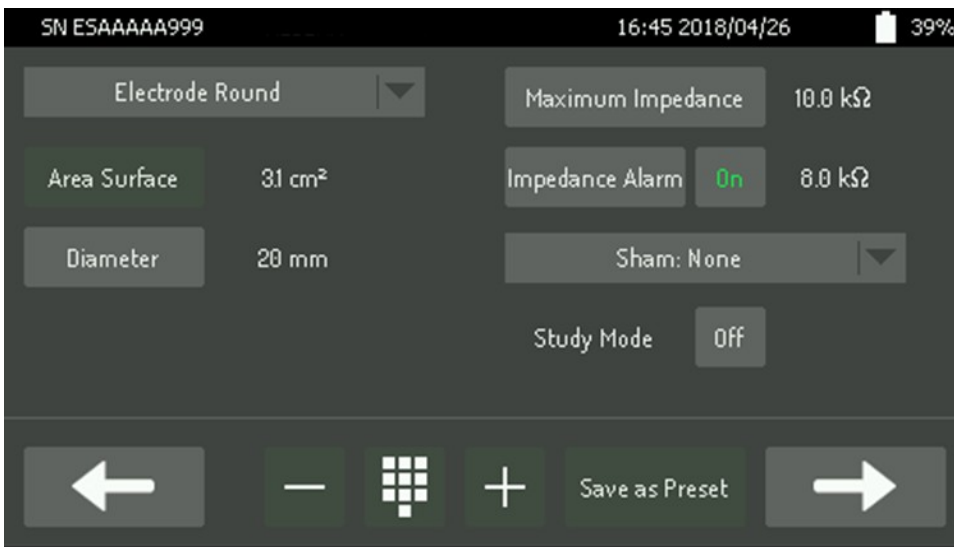


Figure 42. Electrode Round.

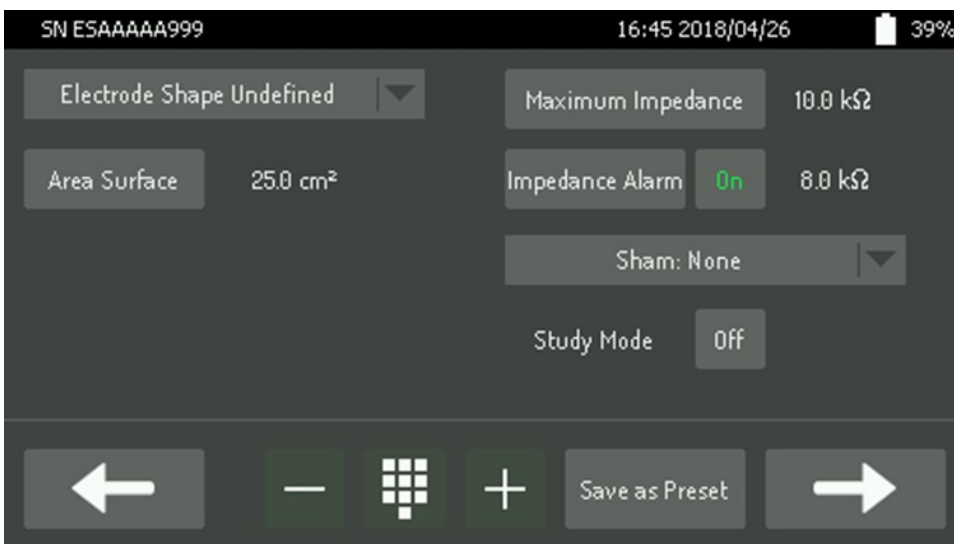


Figure 43. Electrode Shape Undefined.

The electrode shape can be left as undefined, but the surface area of the electrode must still be entered (Figure 43).

When selecting a rectangular electrode shape, enter two dimensions (a and b) to change surface area. Once selected the letter in the box will change to green Use + and – to change dimensions by 5mm (min. 5mm, max. 300mm). Pressing the keypad icon will allow the user to enter the exact number (Figure 41). The area is calculated automatically.

To edit the surface area of round electrodes it is necessary to enter the diameter. Use + and – change dimensions by 5mm (min. 12mm, max. 300mm). The keyboard may also be used (Figure 42).

To edit the surface area of an undefined electrode shape press 'Area Surface'. Use + and – change dimensions by 1cm² (min. 1cm², max. 900cm²). The keyboard may also be used (Figure 43).

Should a value be entered that exceeds the permitted maximum, it will be automatically corrected to the highest permissible value. If a maximum number of digits is entered, the numeric keypad is deactivated.

Selecting the 'ignore size of electrode' option means that the device will not check the parameter's propriety. For example, using too big current for small electrodes may cause burn or electric shock to the subject.



NOTE! Before choosing the “ignore electrode size” option check if the set parameters will not harm the subject.

‘Ok’ is available when an appropriate value is entered.

‘Back’ closes the keypad and returns to the previous screen without saving.

‘Clear’ erases the entered value so that a new value may be entered.

4.11.2 Maximum Impedance

This option relates to the maximum permissible impedance between two electrodes. Should the impedance exceed the entered value, the stimulator signals an error and terminates the stimulation.



Note: The stimulation can still be terminated or prevented from starting if the current level and impedance cause an overvoltage error, even if the maximum impedance level has not yet been reached.

To edit the impedance level, select ‘Maximum impedance’ from the advanced options menu. Use + and – to increase/decrease the level in increments of 1k Ω . Pressing the keypad icon will bring up the keypad screen (Figure 44).

The valid range of maximum impedance is from 10k Ω to 50k Ω . To enter fractional values, press the decimal key on the keypad. If an entered value exceeds the permitted maximum, it will be automatically corrected to the highest permissible value. If two digits are entered for the non-fractional part, the numeric keypad will be deactivated, however you can continue to enter decimal places, by pressing the decimal key (Figure 45). When the total number of digits is entered, numeric keypad will be deactivated.

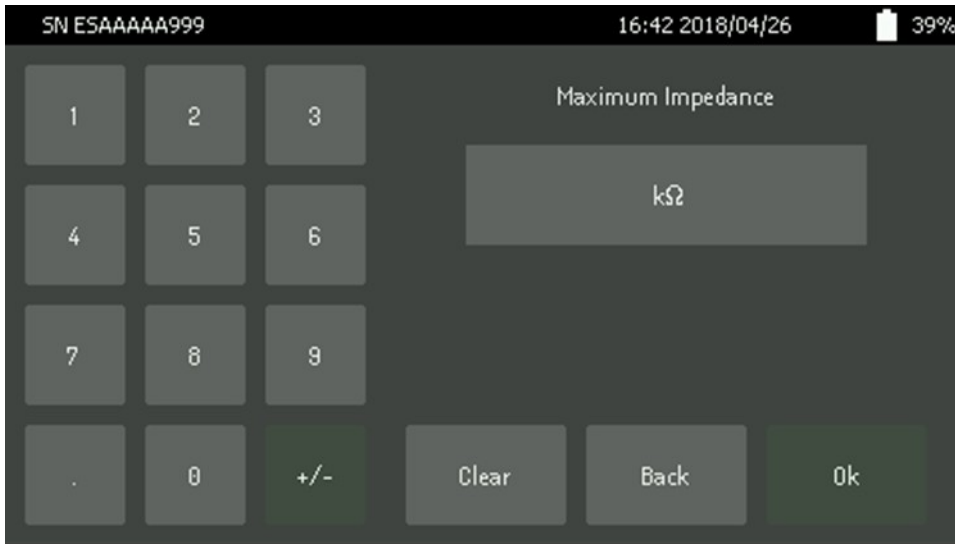


Figure 44. Maximum impedance setting.



Figure 45. Maximum number of digits entered for maximum impedance.

'Ok' is available when an appropriate value is entered.

'Back' closes the keypad and returns to the previous screen without saving.

'Clear' erases the entered value so that a new value may be entered.

4.11.3 Impedance Alarm

An impedance alarm will sound if the impedance exceeds the value entered here. Stimulation will not be interrupted.

This parameter is optional. It can be enabled/disabled by selecting on or off on the advanced options screen. If the on/off button displays green 'on', the alarm is enabled (Figure 46). It can be edited in a similar manner as the maximum impedance parameter. The acceptable range is 5kΩ to 20kΩ.

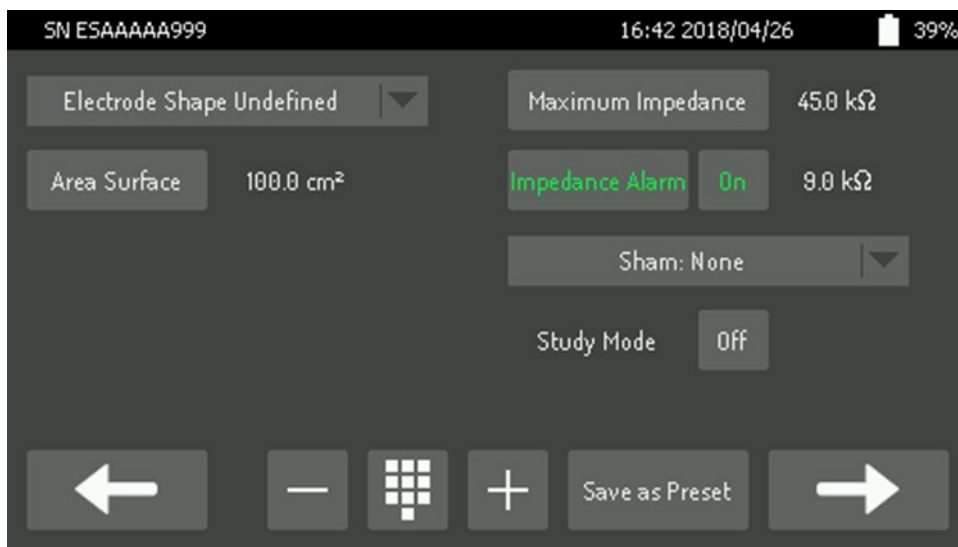


Figure 46: Impedance alarm text showing "on" in green. Alarm has been enabled with user set value shown to the right.

4.11.4 Sham Mode

The sham feature allows the user to determine whether genuine or faux stimulation is administered. Three options are available to select in the drop down.

- None – Normal stimulation (Figure 47).
- Single Blind – Always sham stimulation. The recipient does not know whether stimulation is occurring or not, but the researcher does (Figure 48).
- Double Blind – normal or sham stimulation determined by a PIN code (Figure 49). Double Blind mode maintains complete reliability of the research - the subject does not know what kind of stimulation and what stimulation parameters are administered. The researcher/operator, in turn, knows the settings, but does not know whether the proper stimulation or placebo stimulation will be given. Only the research manager decides what the study scenario is: whether it is a proper stimulation or a placebo one.



Figure 47: Sham: None

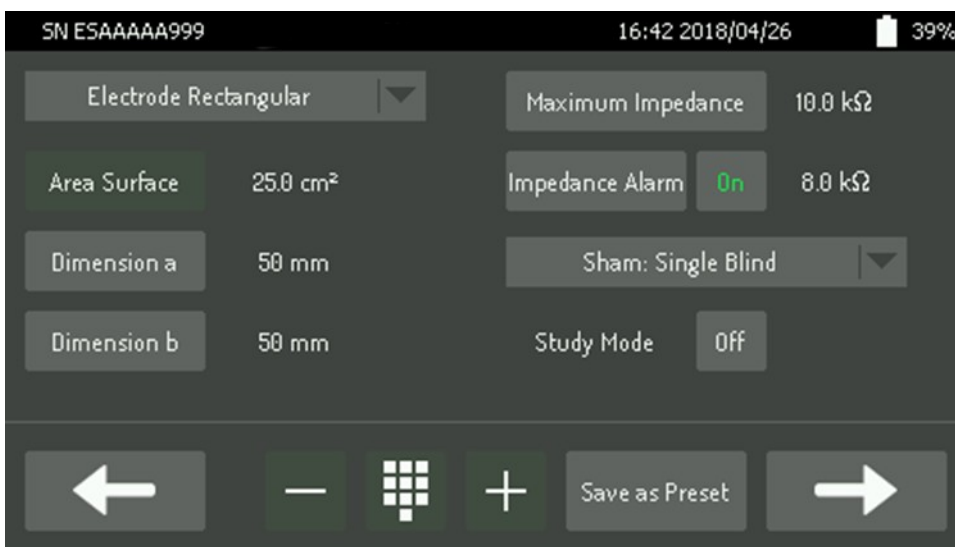


Figure 48: Sham: Single Blind.

During the Double Blind sham stimulation some of the parameters will not be displayed or will be displayed in an obscured way.

The operator only enters an appropriate PIN code (shared by the research manager) to start the stimulation (Figure 50). Some PIN codes trigger normal stimulation and others trigger the placebo one. A list of 30 five-digit PIN codes needed to initiate Double Blind is included with this manual. For more PIN codes, please refer to the distributor.



Figure 49: Sham: Double Blind.

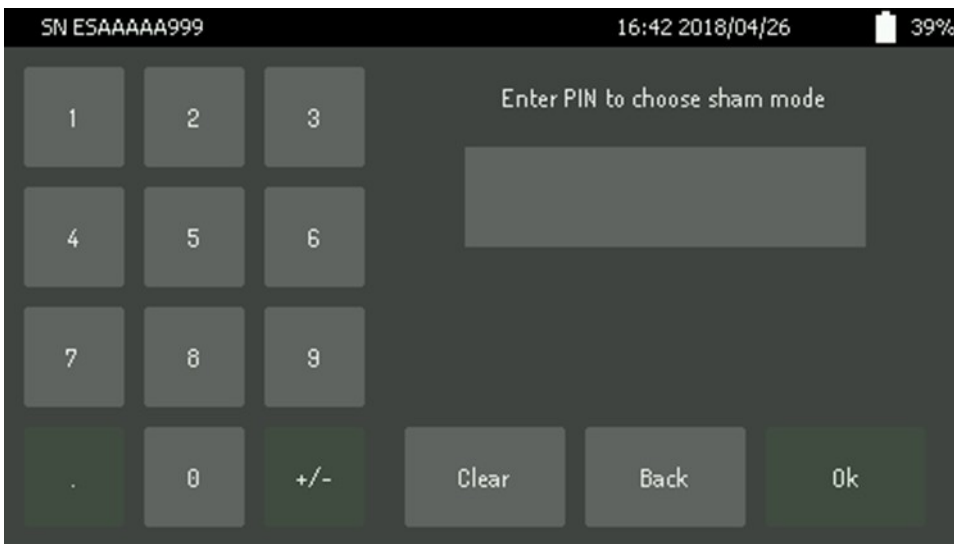


Figure 50: Enter PIN to choose sham mode.

During stimulation the operator can check whether sham stimulation is turned on or not by polling stimulation status via SCPI protocol. To set a sham option, select from the sham drop down menu. Information about sham mode and eventually given PIN code are stored in log files on the device (if enabled) and can be used for further post processing of data.

4.11.5 Study Mode

Study mode is an advanced option that hides the stimulation parameter details during stimulation. When study mode is active, there are no parameters such as amplitude or frequency displayed on the Summary screen (see chapter 4.14) – only impedance check is visible for safety reasons (Figure 51).

Impedance display in Study Mode is limited to colour-code, displaying a red bar when maximum impedance is exceeded, yellow bar when impedance alarm is exceeded and green bar when neither is exceeded. Numerical value is hidden on purpose.

Study mode is used in stimulation situations where the subject must not see the stimulation parameters so as not to influence the results of the experiment.

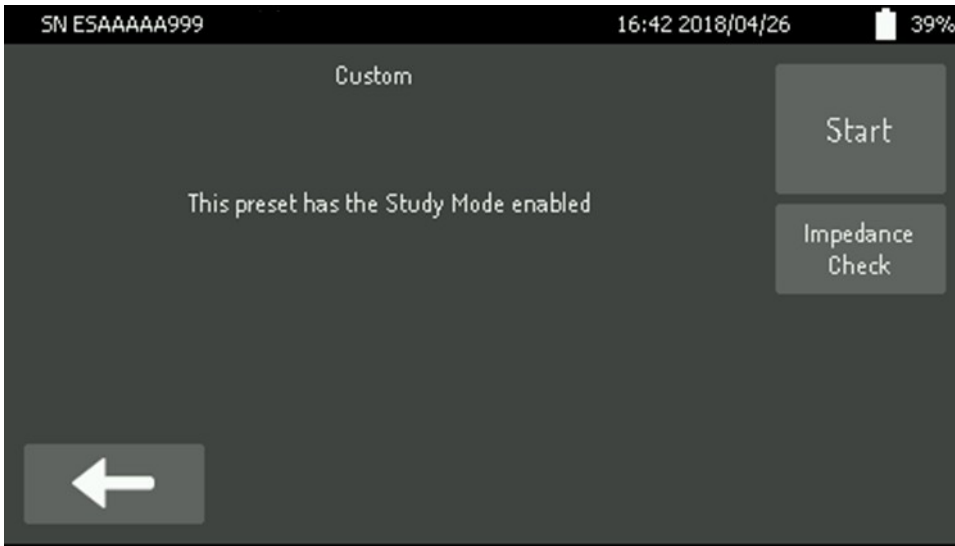


Figure 51: Study mode summary screen.

4.11.6 Test Mode

Test mode allows for changing the current amplitude during ongoing stimulation without the need to stop it. This facilitates the selection of appropriate parameters, especially when the patient reports discomfort. Test mode is available both through the user interface (Figure 52), and via the API. To activate it, set the switch to the "on" position and start the stimulation.

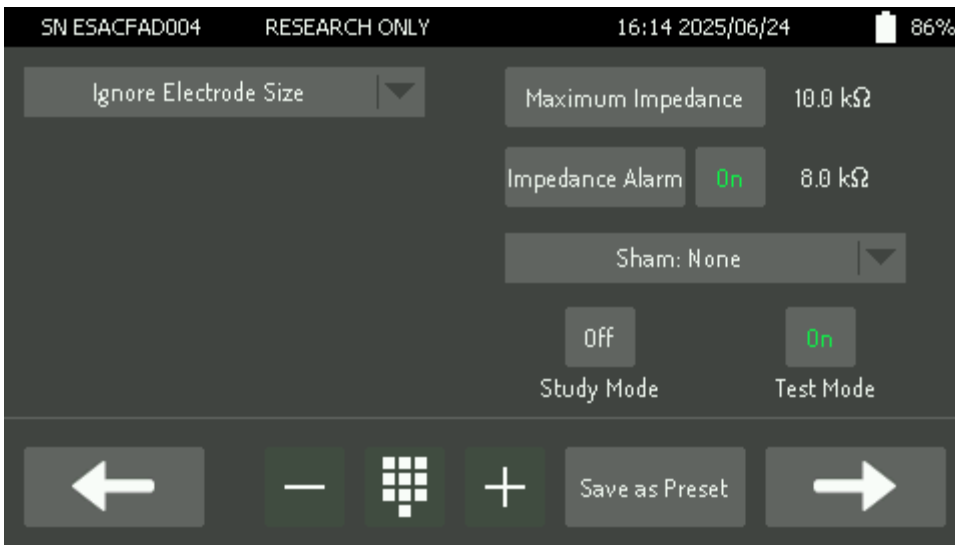


Figure 52. Turning on test mode stimulation.

The screen features "+" and "-" buttons that allow adjusting the current amplitude in 100 μ A increments (Figure 53). After stopping the stimulation, the new values are retained and can be saved in a predefined protocol. However, it should be noted that enabling test mode itself is not saved.

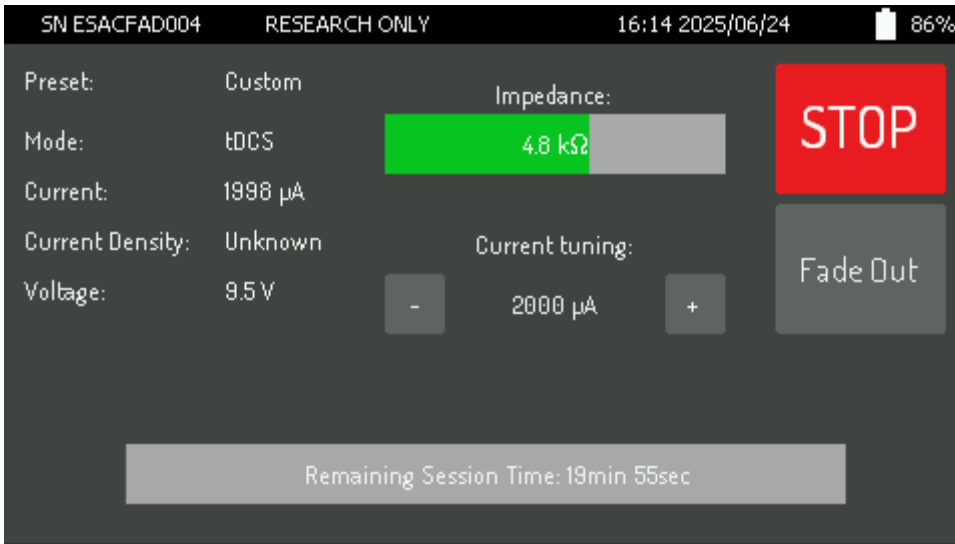


Figure 53. Stimulation running in test mode. The + and - button allows to change current intensity in real time.

4.12 Saving Presets

Once all of the above parameters have been selected, the user is able to save them as a preset which can be selected in first screen (Figure 10). To do this, press 'Save as Preset'. This will direct the user to the presets window (Figure 54).



Figure 54. Presets screen with slot selected for save.

To save a preset, select any of the buttons which show a preset slot. The selected slot will then be highlighted green and the 'Save' button in the bottom right corner will become available (Figure 54).

When the preset is saved, it will be stored in the chosen slot. If the selected slot is not empty, any previously saved preset data will be overwritten. Presets saved this way will be given a generic name ("Preset_4" for example). It is not possible to change the name of these presets via the touch screen.

Saving a preset and giving it a custom name can only be done via the USB interface using the serial terminal and SCPI protocol or via the device computer application. Connecting via SCPI protocol is described in the Appendix to this manual. Deleting stored presets must also be carried out via SCPI protocol.

After setting all the stimulation parameters in the Custom Stimulation window, you can save the stimulation prepared in such a way as a preset, which will then be visible in the "Presets window" of the main menu.



A preset cannot be saved if the maximum safe current density is exceeded. When this is the case, the Save button will not be illuminated. Whether or not stimulation is possible is shown in the Summary window.

To exit the presets screen, press the arrow in the bottom left corner.

4.13 Limit Mode

It is possible to restrict the functionality of the device to only load presets that had been saved earlier. In this mode the parameters of stimulation and electrode cannot be adjusted and it is only possible to invoke a preset without any alterations.

Switching limit mode on or off is only possible by connecting to computer by USB cable and using Corstim application (or low-level terminal), which is outside the scope of this manual. Switching the mode on and off is PIN-code protected.

When the device is in restricted mode, it automatically opens the predefined protocol selection screen upon startup. To start stimulation, select the desired preset, which will be highlighted in green, and press the Load button (Figure 55).

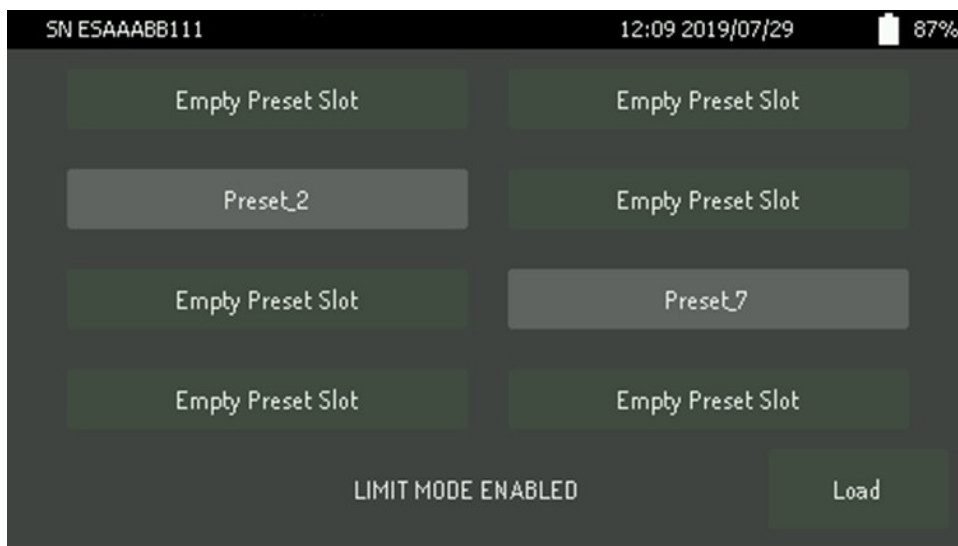


Figure 55. Preset selection as main screen in limit mode.

4.14 Summary

Once all of the parameters have been set and the user is ready to proceed to stimulation, press the arrow in the bottom right of the advanced options screen (Figure 56).

Before starting stimulation, a screen summarizing the parameters to be administered will be shown (Figure 57). If the maximum safe current density is exceeded, the 'Start stimulation' button will be unavailable, and the parameter will be highlighted in red (Figure 58). If the electrode size was not entered whilst setting the advanced options (the ignore electrode size option was chosen), a warning will be shown (Figure 59) but stimulation can still be started.

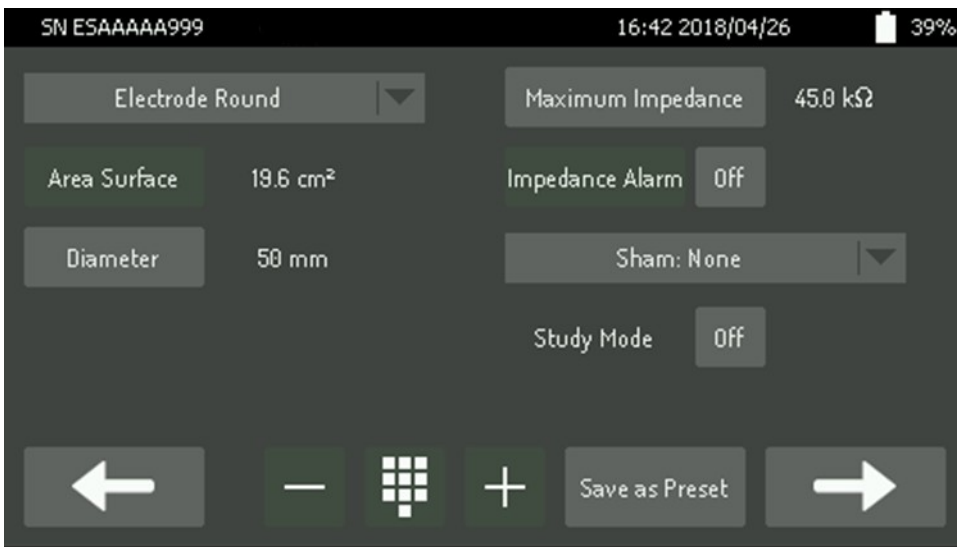


Figure 56. Advanced options screen. All parameters have been set so stimulation can proceed via the right-hand arrow.

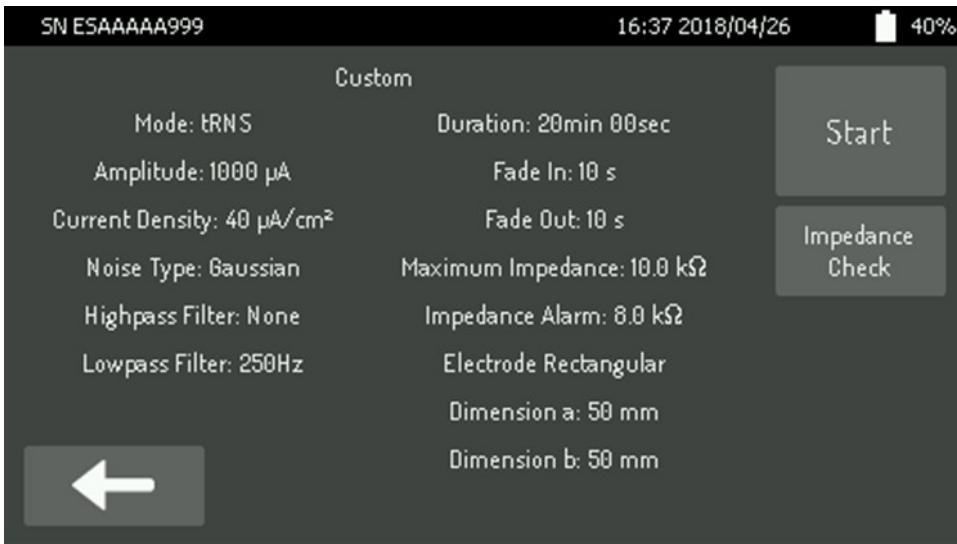


Figure 57. Summary screen showing stimulation settings.

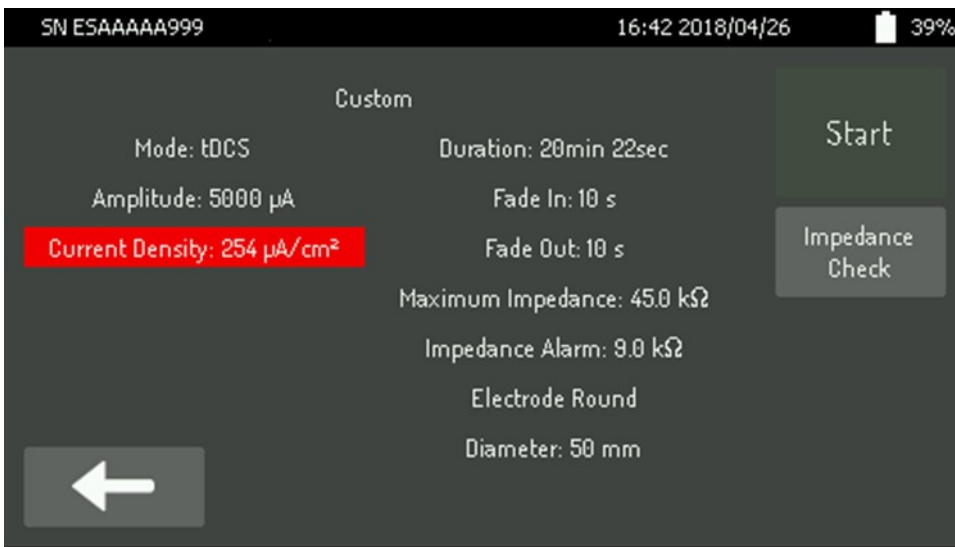


Figure 58. Current density is too high, so stimulation cannot proceed.



Figure 59. Current density is unknown as electrode size has not been entered. Stimulation can still start.

4.15 Impedance Check

Before starting stimulation, it is recommended that a manual impedance check is carried out. This will ensure that electrodes are mounted correctly, the required stimulation current can be reached, and overvoltage cannot happen. Pressing 'Impedance Check' button will direct the user to the impedance check screen (Figure 60) and a continuous impedance measurement will start.

If the level of impedance is low, which is a correct and desirable value, a green bar will be shown.

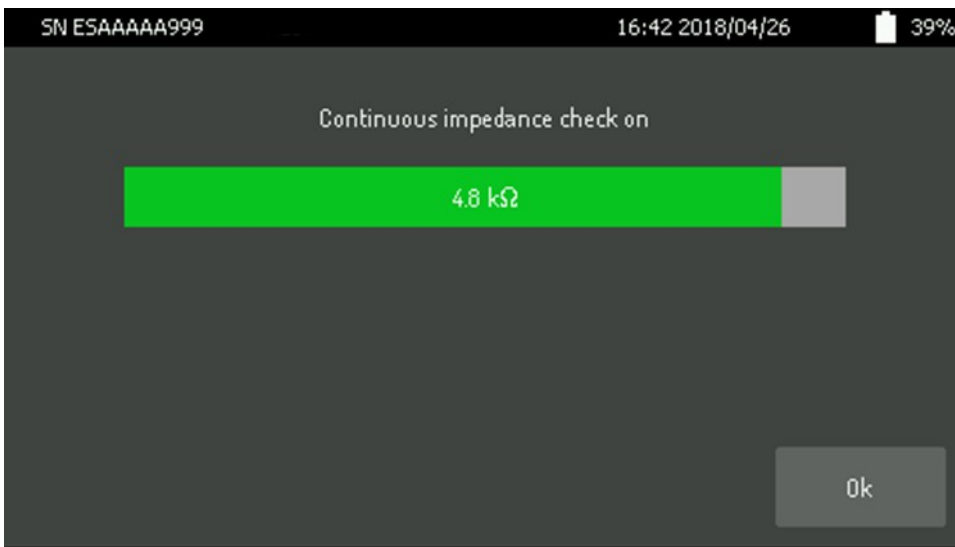


Figure 60. Impedance check screen. Green bar shows a low level of impedance.

If the impedance level exceeds the user defined threshold, the impedance bar will be yellow, and a warning message will be displayed (Figure 61).

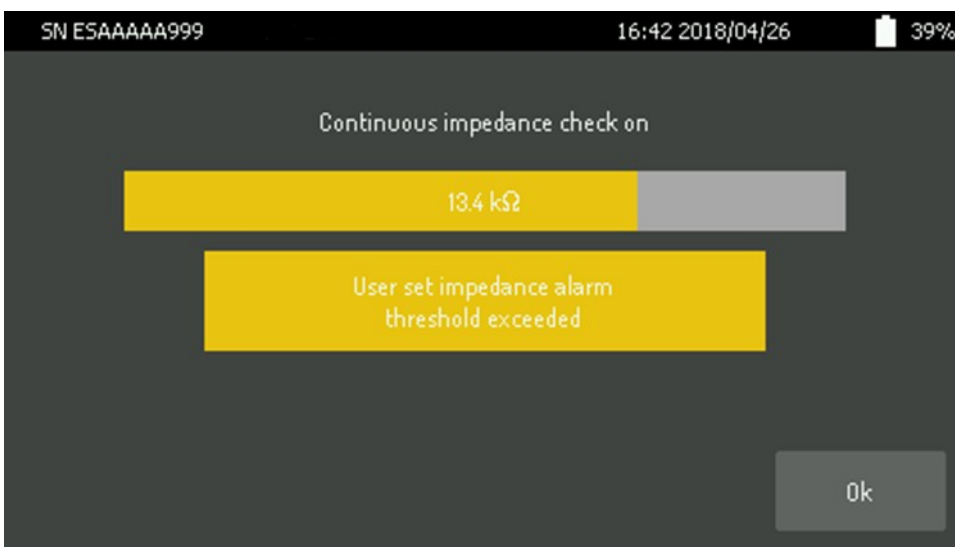


Figure 61. User set impedance level has been reached. Impedance bar is yellow, and a warning message is shown.

If the measured level of impedance is so high as to cause overvoltage, the impedance bar will be red, and a second warning will be shown (Figure 62).

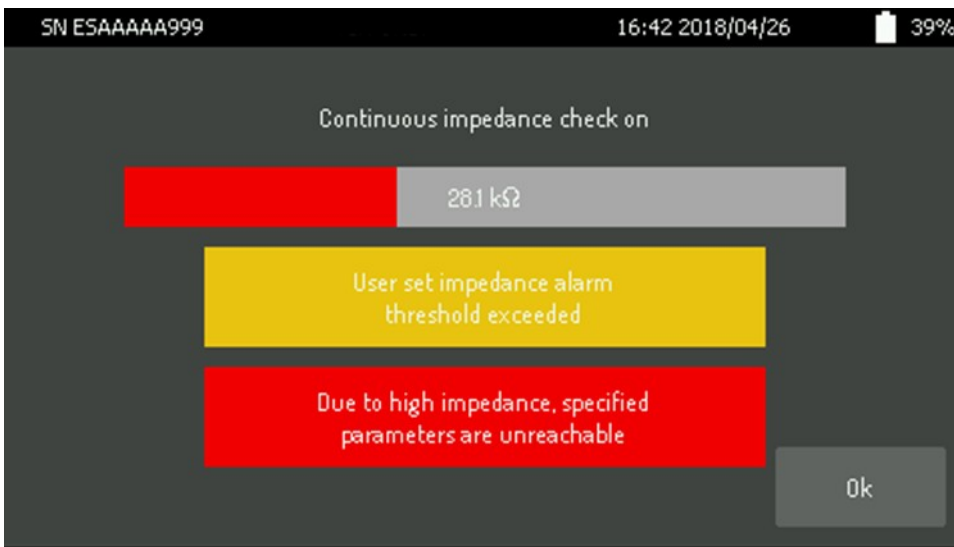


Figure 62. Specified parameters would not be achievable, so secondary warning message is displayed in red.

If the measured impedance exceeds the maximum level, both warning messages will be shown in red (Figure 63).

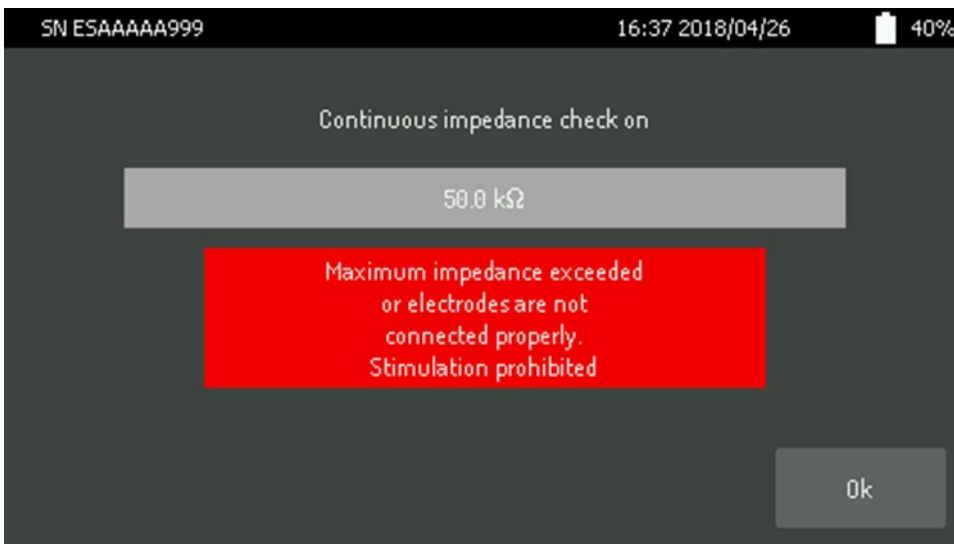


Figure 63. Maximum level of impedance reached. Both warning messages show in red.

If the impedance level measured is too low or electrodes are shorted, the impedance bar will be shown in red and a dedicated warning message will be shown in red (Figure 64).

The measured impedance value during impedance check and during proper stimulation can differ due to the characteristics of the measuring signal.

If impedance level is deemed to be an acceptable level, press the 'Ok' button in the bottom right corner to return to the preset summary screen. To start stimulation, press 'Start Stimulation'.

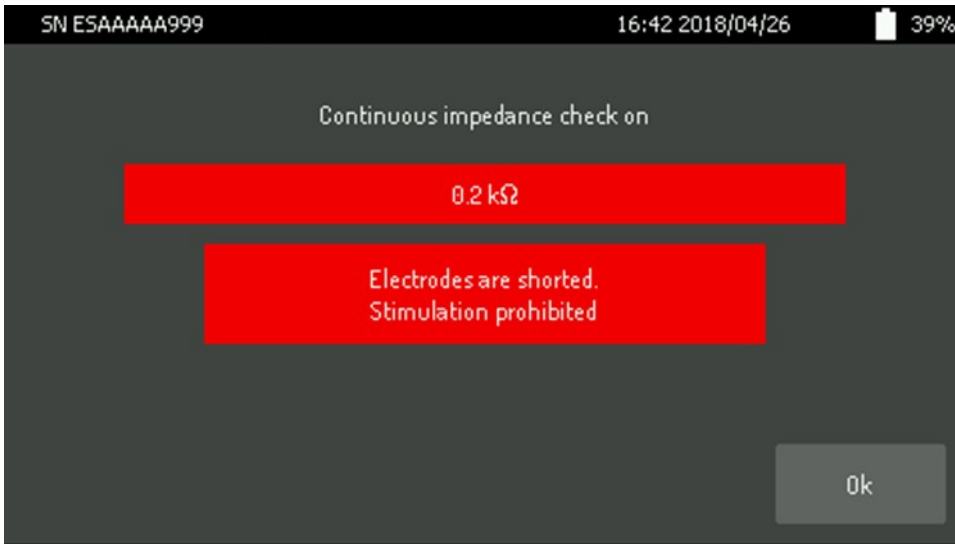


Figure 64. Short circuit detected - impedance bar shown in red with a specific warning message.

4.16 Stimulation

Stimulation starts with a single impedance check. If it is within accepted levels, actual stimulation begins. This feature can be turned off via SCPI protocol, but for safety purposes, each dose of stimulation requires an impedance check.

The stimulation screen shows current preset name, measured values, user set values where applicable (frequency and duty cycle, for example), progress bar and stop buttons. During Fade in/Fade out the measured values are unavailable (Figure 65). Impedance bar shows colour-coded indication of impedance with no numeric value.



Figure 65. Stimulation screen showing mode, impedance bar. Measured values are shown as "Fading in" or "Fading out" during the fade in and fade out periods.

During normal stimulation, all measured parameters are displayed (Figure 66). In some cases (like very low tACS frequencies) parameters are shown as 0 until first valid measurements are gathered. In 0.01Hz tACS this could take up to 50 seconds.

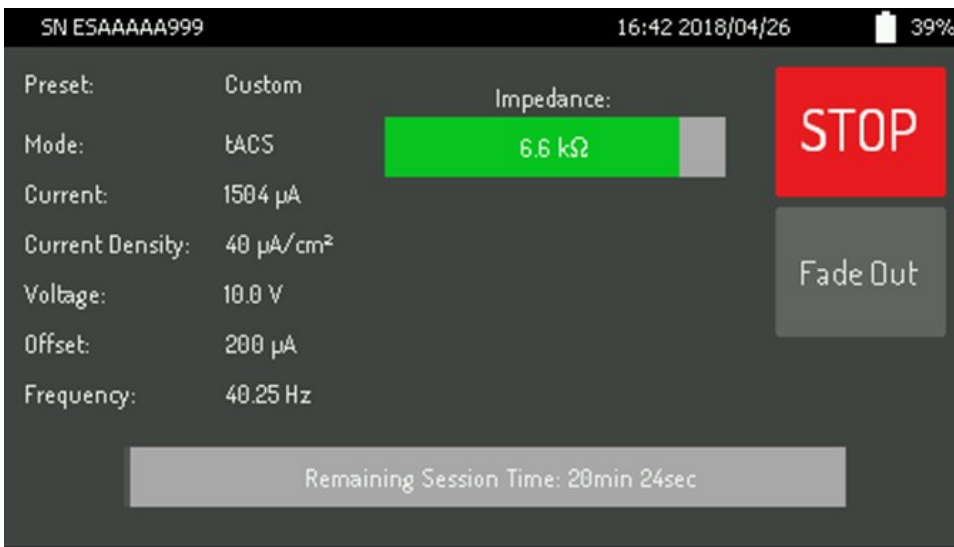


Figure 66. Stimulation screen during normal stimulation. All parameters are displayed.

If the impedance alarm has been set, which is triggered during stimulation or impedance becomes too high or low, the impedance bar will turn the relevant colour and the corresponding warning message will be shown, as seen in the impedance check screen in 4.15 and Figure 67.

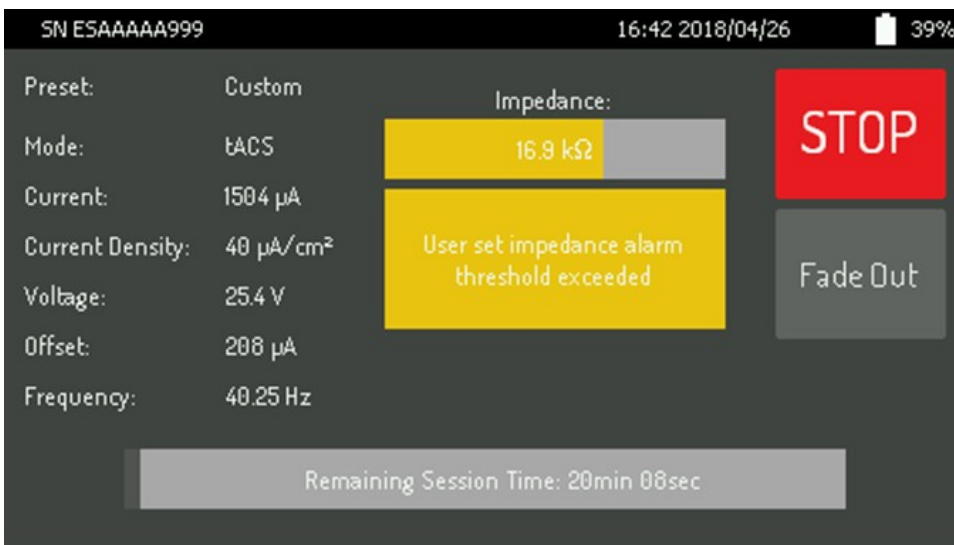


Figure 67. Impedance warning shown during stimulation. Stimulation will still be administered at this point.

If the measured impedance levels become too high, regardless of the set user limit, stimulation will be aborted, and the emergency fade out will begin (Figure 68).

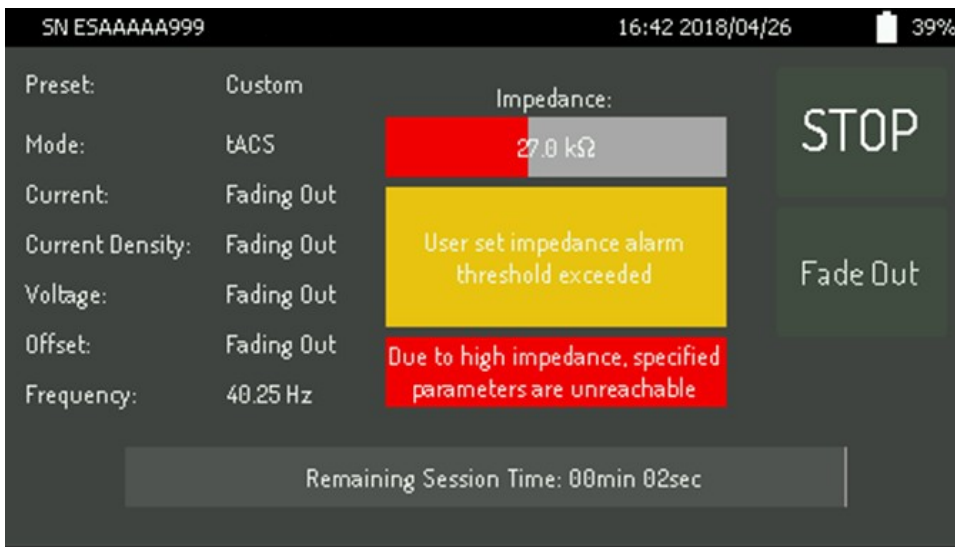


Figure 68. Stimulation has been aborted due to high impedance levels. Emergency fade out is underway.

To stop stimulation manually, press either 'STOP' or 'Fade out'. Pressing 'Fade out' will cause stimulation to cease and proceed to a pre-defined fade out length selected in the stimulation settings screen and then stop.

The 'STOP' button initiates an emergency fade out (2 seconds) and should only be used in an emergency, when it is necessary to stop the stimulation as fast as possible. The use of the 'Fade out' button is recommended so as not to cause discomfort to the subject.

The stimulation will not start if:

- parameters are unreachable (Figure 69),
- impedance measurement is outdated (Figure 70),
- battery is low (Figure 71).

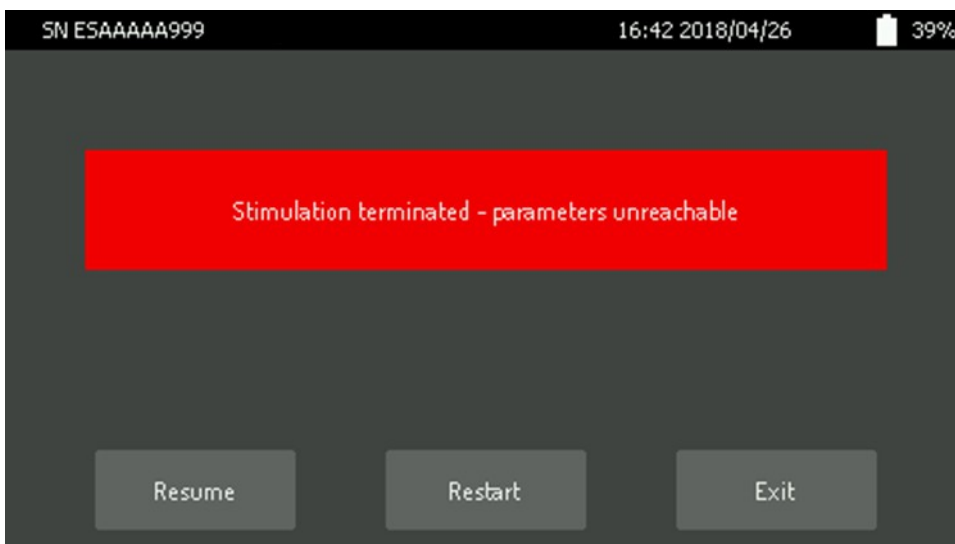


Figure 69. Stimulation terminated - parameters unreachable.

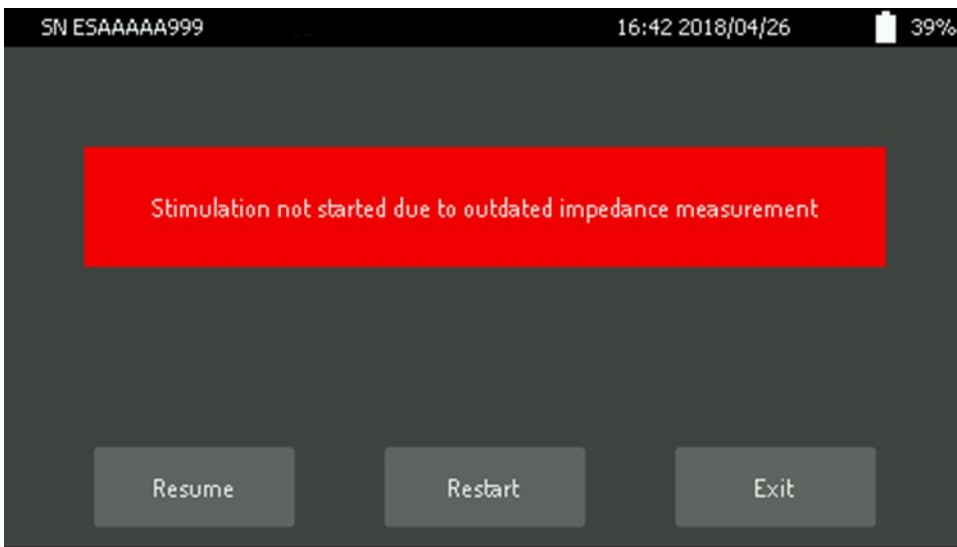


Figure 70. Stimulation not started due to outdated impedance measurement.

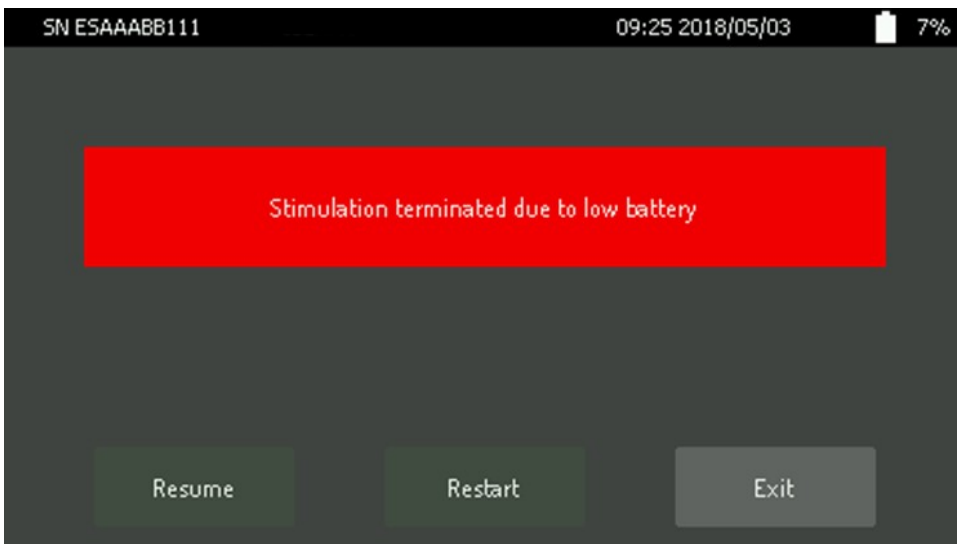


Figure 71. Stimulation terminated due to low battery.

4.17 End of Stimulation

After a successful stimulation dose, the exit screen is shown with a message confirming the end of stimulation (Figure 72).

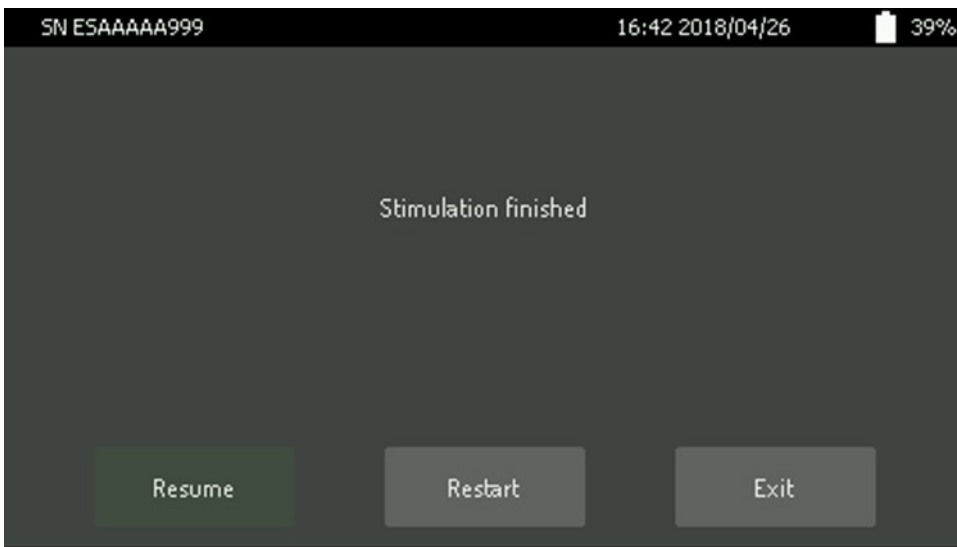


Figure 72. Screen showing that stimulation has finished without errors.

If stimulation was terminated automatically (by a warning) or early (manually) the appropriate alarm message is show in the exit screen.

If a stimulation has ended prematurely it can be resumed (if it is possible to stimulate) by pressing 'Resume' which opens a preset summary screen (Figure 57). However, stimulation time will be decreased by the amount of time that elapsed in the aborted session (Fade in and Fade out are not counted as stimulation time).

The stimulation will finish automatically if:

- the programmed stimulation was successful (Figure 72),
- impedance value falls below or above safe range (Figure 73),
- user connects a power unit to the device (Figure 74),
- user disconnects electrodes or pin cables (Figure 73),
- user presses 'STOP' (Figure 78),
- user presses 'Fade out' (Figure 78).

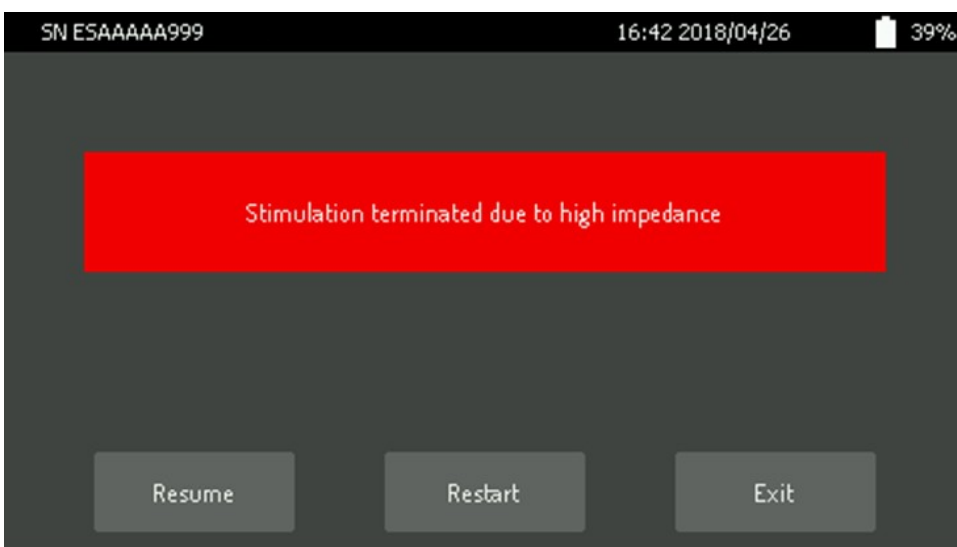


Figure 73. Stimulation terminated due to high impedance or disconnected electrodes – visible as high impedance.

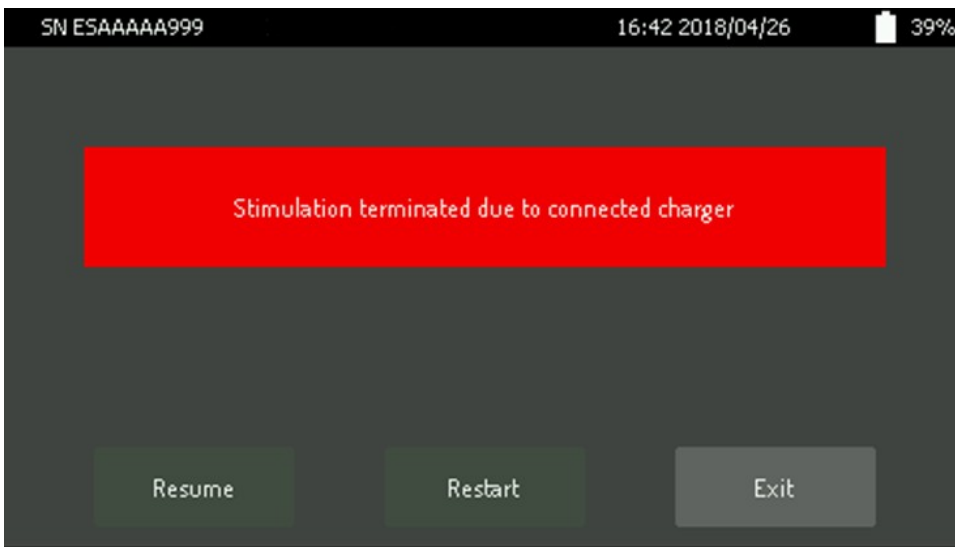


Figure 74. Stimulation terminated due to connected charger.

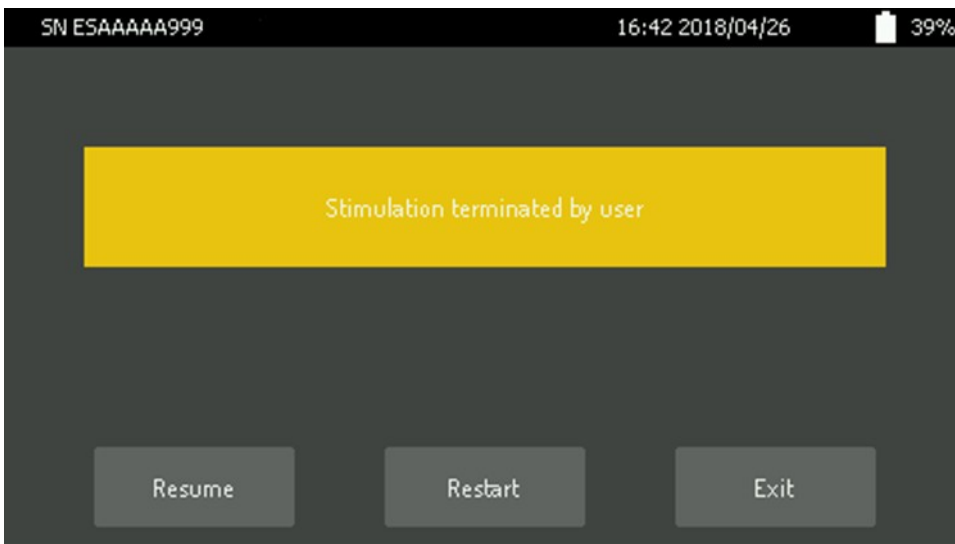


Figure 75. Stimulation terminated by user.

If the reason for the aborted stimulation was due to an impedance error, a new impedance check will have to be carried out. This should be completed before going to the preset summary screen.

Stimulation can be restarted (if it is possible to stimulate) by pressing the 'Restart' button. This will take the user to a preset summary screen (Figure 57). Stimulation time will have been reset to the original value established in the stimulation parameters.

'Exit' takes the user back to the home screen (Figure 10).

4.18 Loading a Previously Saved Preset

To load a stimulation protocol from a preset, select 'Presets' from the home screen (Figure 10)

Stored presets will be shown as available buttons with preset names. Select the preset to be used. The chosen preset will then be highlighted green and the 'Load' button will become available (Figure 76).



Figure 76. Preset selection screen. Preset 2 chosen for loading.

Once 'Load' has been pressed, the user is taken to a new summary screen, showing the parameters about to be administered. Note that on the summary screen, Preset 2 is shown as selected (Figure 77). To load a different preset, press the bottom left arrow.



Figure 77. Preset summary screen. Note that Preset_2 has been chosen.

4.19 Analogue Input Mode

Analog input mode is a special mode of operation, in which patient's current is generated in direct proportion to an externally-provided analogue control voltage.

Analog Input Mode can be entered from the device's home screen (Figure 10). The device in this mode operates as follows:

- accepts analogue voltage signal from -2.5V to +2.5V range from an external source, such as signal generator
- generates current stimulation waveform directly proportional to external signal, at conversion factor of 2mA of patient per 1V of external signal voltage, up to -5mA to +5mA range
- provides adequate medical-grade galvanic isolation between external analogue source and patient

In Analog input mode there are no parameters to be set. Once chosen, the stimulation can start or Manual impedance check may be invoked (Figure 78). Maximum impedance accepted in this mode is set to 50k Ω and may not be adjusted. As a safety feature, this stimulation mode can be terminated by open-circuit or overvoltage detection like in other stimulation modes.



Figure 78. Summary screen in Analog Input Mode.

During analog input mode stimulation there's no duration limit and elapsed stimulation time is displayed instead. Additionally, current, voltage and impedance indication is provided (Figure 79). When analog input mode is active and patient current is below 50 μ A, measured impedance result is unavailable.



Figure 79. Stimulation screen in Analog Input Mode.

4.20 Remote Mode

Remote mode enables the device to be controlled via SCPI protocol using USB port. The device is visible by the computer as a serial port. As a low-level alternative it is possible to use any serial port terminal application (although it is recommended to use RealTerm). Connection parameters are irrelevant. Please ensure that STM Virtual COM Port Driver has been installed before connecting the device in this way.

When the device is connected to a computer "PC CTRL" or "STER. PC" depending on the user interface language settings is displayed on the upper bar. If the device is not stimulating, the screen changes to the Preset summary screen with the

last set parameters. Buttons are hidden, and all control is done remotely on the computer (Figure 80). The screen is refreshed with every change to Preset.

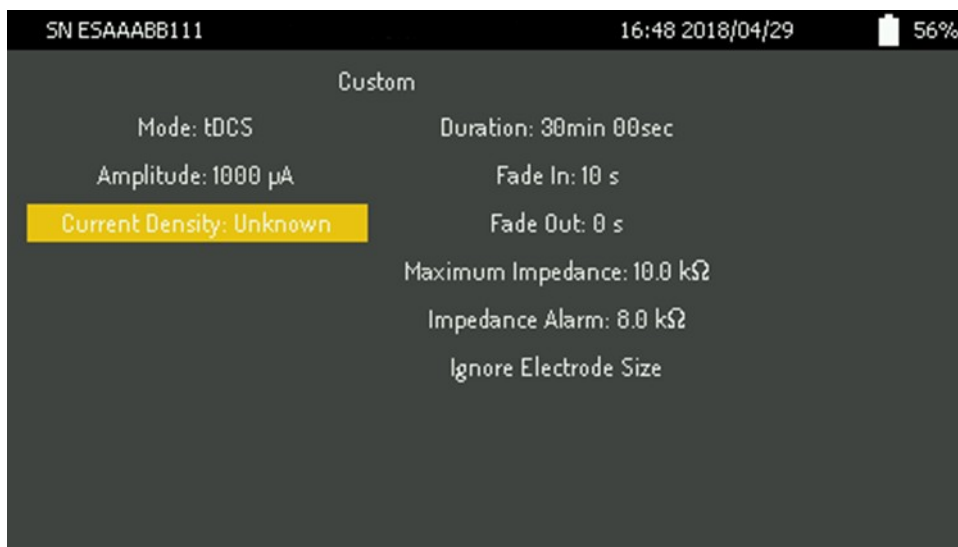


Figure 80. Preset summary screen in Remote Mode.

During stimulation, for safety reasons, 'Stop' and 'Fade Out' buttons are still active (Figure 81). When stimulation ends, the screen reverts back to summary screen.

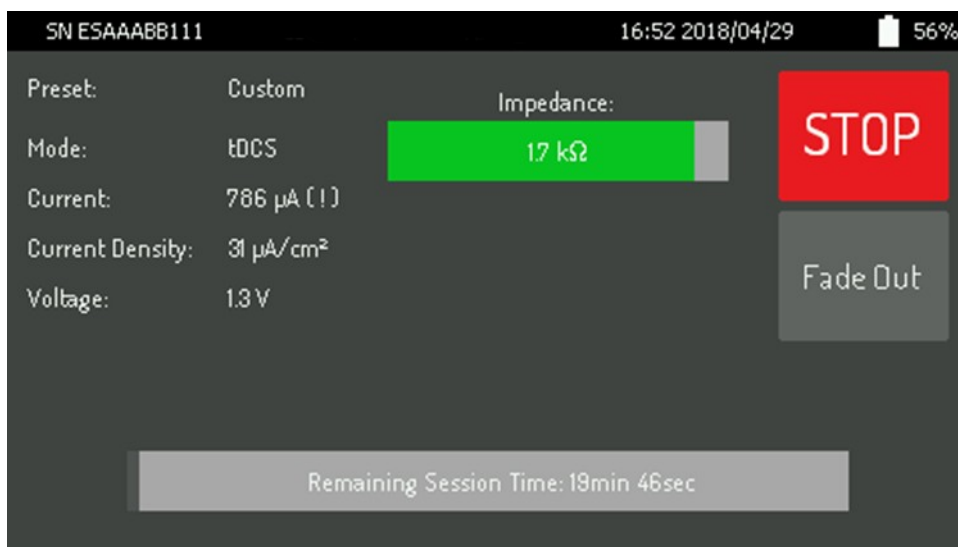


Figure 81. Stimulation screen in Remote Mode.

Remote mode allows full control of all device functions, some of which are not otherwise available from the touch screen user interface (such as Triggers). For a list of all commands, with examples of use, see Appendix to this manual.



It is recommended to use the original dedicated application to connect the device to a computer. Any deviation from this recommendation is done at the user's own risk, and neither the manufacturer nor the distributor can be held responsible for any problems that may result.

Disconnecting the USB cable will cause the device to return to the start screen; however, it is recommended to turn off the device before disconnecting the connection.

5 Triggers and Analogue I/O

Triggers are discrete or digital input and output signals that allow Corstim to synchronise with other devices such as EEG, TMS etc.

Analogue Input signal is used for stimulation governed from external signal source.

Analogue Output signals are used to monitor the patient current and further process or record it.

5.1 Triggering

Input and output control signals (triggers) allow the user to integrate the stimulator with other laboratory devices and use specific events during the stimulation protocol execution to trigger external actions or use external signals to drive the operation of the stimulator. The functionality of input and output triggers can be used via the SCPI protocol as well as the user interface. Trigger settings can be saved as part of a protocol (preset) in the device's memory for easy future use without the need for reconfiguration.

5.2 Analogue Input

Analogue input signal is used along Analogue Input Mode stimulation. In this mode patient stimulation current is closely following the voltage waveform at the Analogue Input provided from the external source. The transfer ratio is non-adjustable at 2mA of stimulation current per 1 volt at analogue input.

The waveform presented to the analogue input may be arbitrary (AC, DC, noise etc.) as long as the signal stays within $\pm 2.5V$ range and is bandwidth limited to 1kHz (at -3dB attenuation).

'Fast' signals near 1kHz and above will be transferred with considerable attenuation, with visible slope and/or phase shift.

Analogue Input Mode cannot be used simultaneously with the Remote Mode. Neither application nor SCPI commands allow to manage Analogue Input Mode.

Corstim provides measurements in of output current, voltage and estimated impedance, thus ensuring to control skin-electrode contact. Indications of measurements in Analogue Input Mode are based on root-mean-square (RMS) values averaged over a period of time, thus indicated values are always positive values and have definitive relations to other parameters, such as factor of 0.707 to peak value in case of a pure sine wave or equal to absolute DC value in case of DC input.

5.3 Analogue Output of 500mV per mA

Analogue output is a single-ended directly proportional representative of patient stimulation current.

This output is active at every mode of stimulation (TDCS, TACS, TRNS, Analogue input).

Output level is typically convenient for oscilloscope examination of patient current.

Output transfer is also complementary to Analogue Input, therefore allowing for different configurations of multiple Corstim device in multichannel stimulation (mirroring current, daisy-chaining, parallel operation etc.)

5.4 Analogue Outputs of 2.5mV per mA

Low level Analogue Outputs are differential scaled-down outputs. The level and differential format make the signal convenient for usage with EEG amplifier or similar processing.

5.5 Pin Layout

The diagram below shows the layout of the pins in the trigger port and their uses.

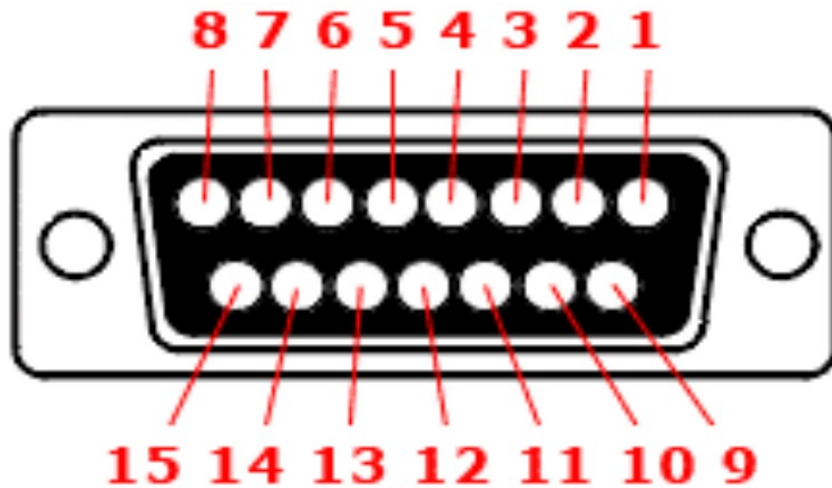


Figure 82. Trigger socket layout.

Pin 1 – Analogue Input	Pin 10 – Analogue Output 500mV/mA
Pin 2 – GND - Ground for triggers	Pin 11 – GND - Ground for triggers
Pin 3 – Analogue Output 2.5mV/mA Negative	Pin 12 – GND - Ground for triggers
Pin 4 – Analogue Output 2.5mV/mA Positive	Pin 13 – GND - Ground for triggers
Pin 5 – GND - Ground for triggers	Pin 14 – Trigger In channel 2
Pin 6 – Trigger Out channel 3	Pin 15 – Trigger In channel 1
Pin 7 – Trigger Out channel 2	
Pin 8 – Trigger Out channel 1	
Pin 9 – 5V Source (max short circuit current: 30mA)	

5.6 Trigger In

Trigger in channels are used to start and stop a stimulation protocol via an external CMOS level compatible source (TTL or 5V CMOS compatible signals). Each trigger channel is configurable independently and works independently. Each channel can be configured so that it can either start or stop the stimulation or do both.

To start the stimulation with a trigger the 'start trigger event' must be configured and a 'start stimulation command' must be sent to the device. The device will then display a 'Stimulation will start after trigger' message and once a trigger event occurs the stimulation will start (Figure 83).

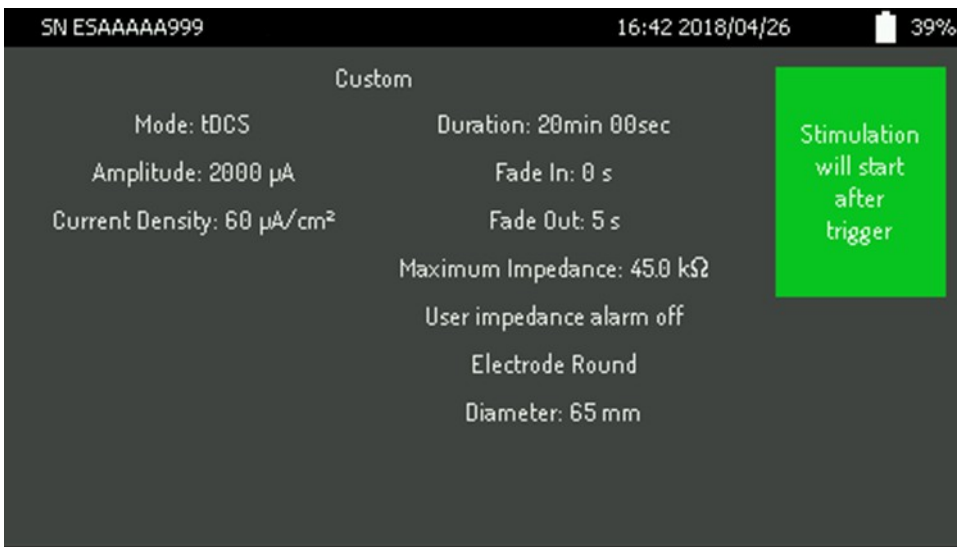


Figure 83. Stimulation will start after trigger.

To stop the stimulation with a trigger the same rules apply: a 'stop trigger event' must be configured and the device must be administering a stimulation.

Each channel can be configured so that it reacts to the following:

- Rising Edge (Figure 84)
- Falling Edge (Figure 85)
- Both Edges (Figure 86)

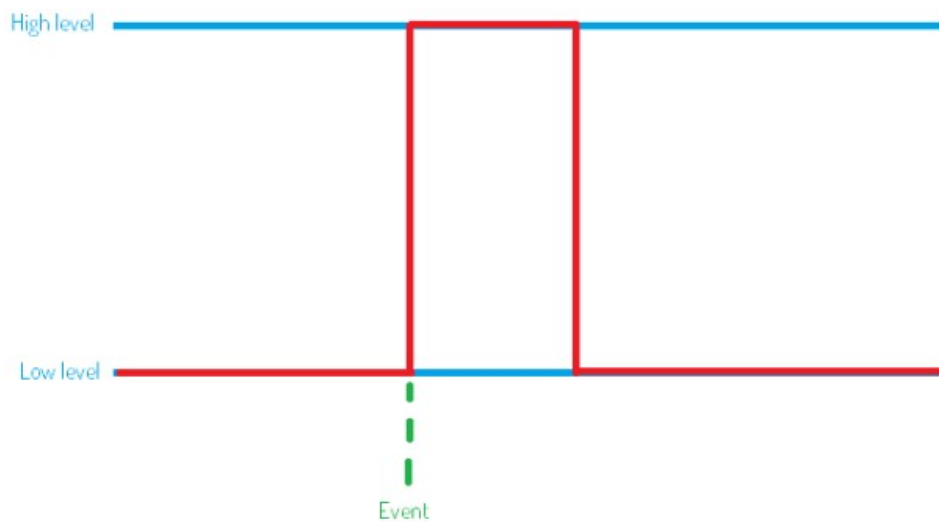


Figure 84. Trigger in setup for rising edge.

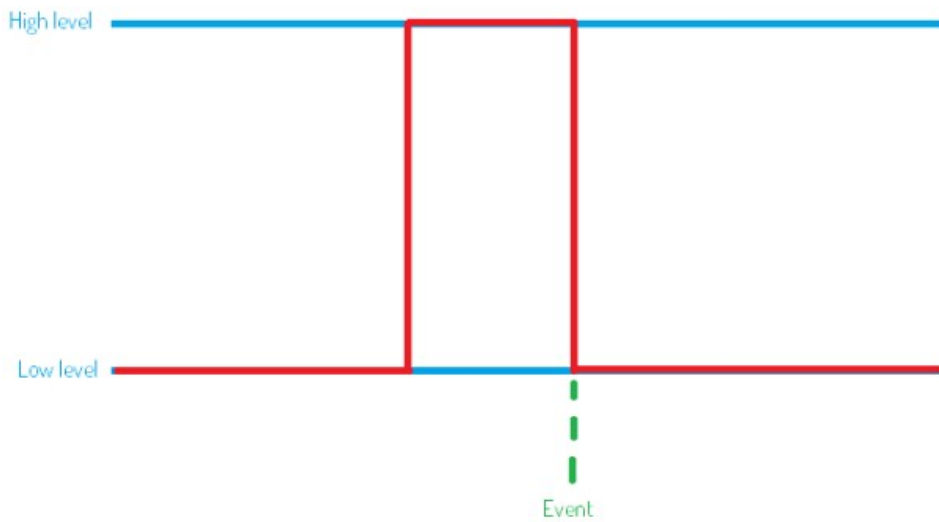


Figure 85. Trigger in setup for falling edge.

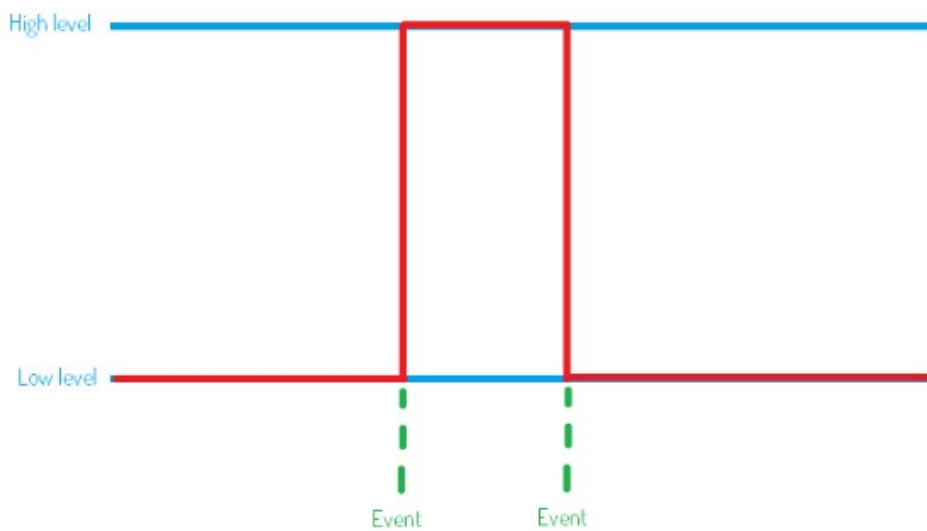


Figure 86. Trigger in setup for both edges.

5.7 Trigger Out

Trigger out channels are used to signal various events during the stimulation to an external device using TTL or 5V CMOS compatible signals. Again, each channel is configurable independently and operates independently. Each channel can be configured with any combination of available events and can be configured to signal events with one of four available modes, such as:

- Pulse High (PH) - Each event is signalled by a signal transition from low to high, then shortly high to low (Figure 87)
- Pulse Low (PL) – Each event is signalled by a signal transition from high to low then shortly low to high (Figure 88)
- Level High (LH) – Only used for signalling the following events: Start or Stop of the stimulation. When stimulation is being administered the signal level is high. When there is no stimulation the signal level is low (Figure 89)
- Level Low (LL) – As above, only used for signalling the following events: Start or Stop of the stimulation. When stimulation is being administered the signal level is low. When there is no stimulation the signal level is high (Figure 90)

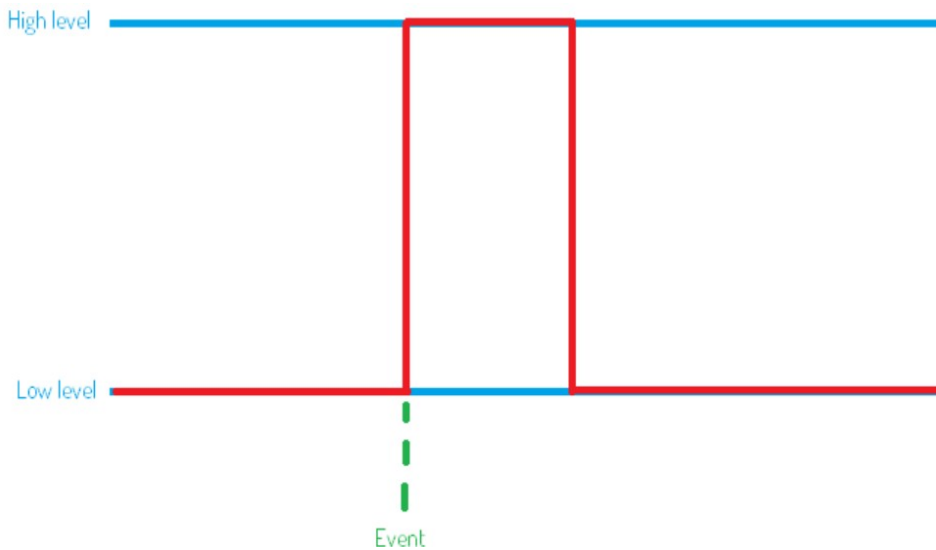


Figure 87. Trigger out setup for pulse high.

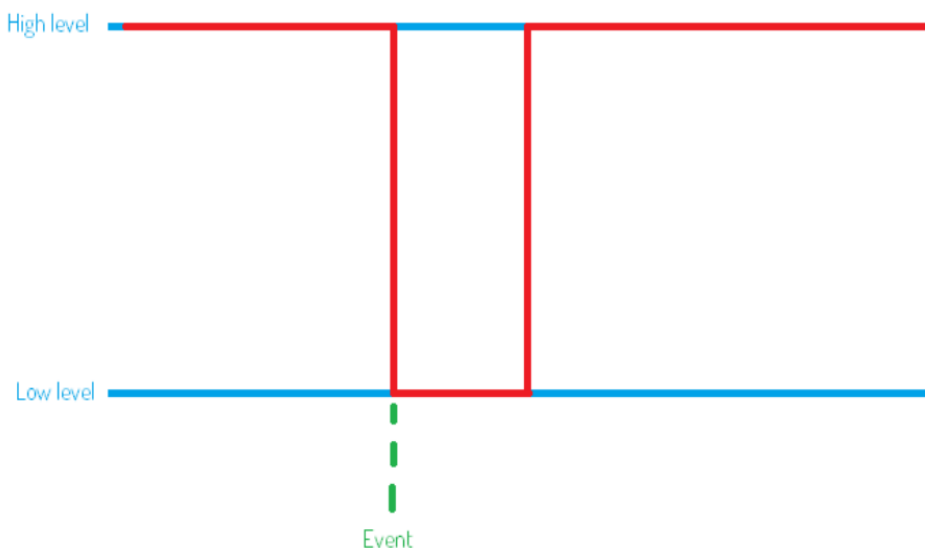


Figure 88. Trigger out setup for pulse low.

Note, that the pulses are very short to accommodate a possible high density of such pulses generated by various events. Any equipment recording such pulses must ensure to have good time resolution not to miss a short pulse. The trigger pulse duration is about ~200 μ s.

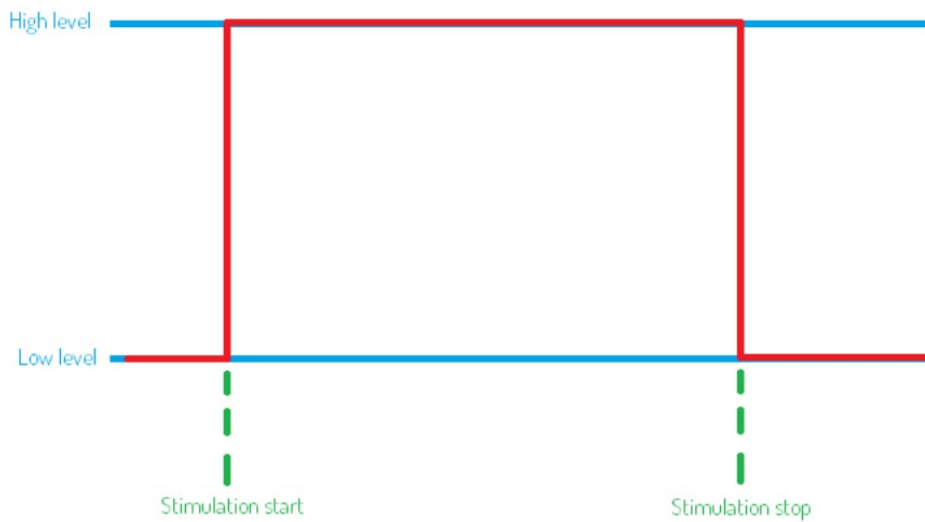


Figure 89. Trigger out setup for level high.

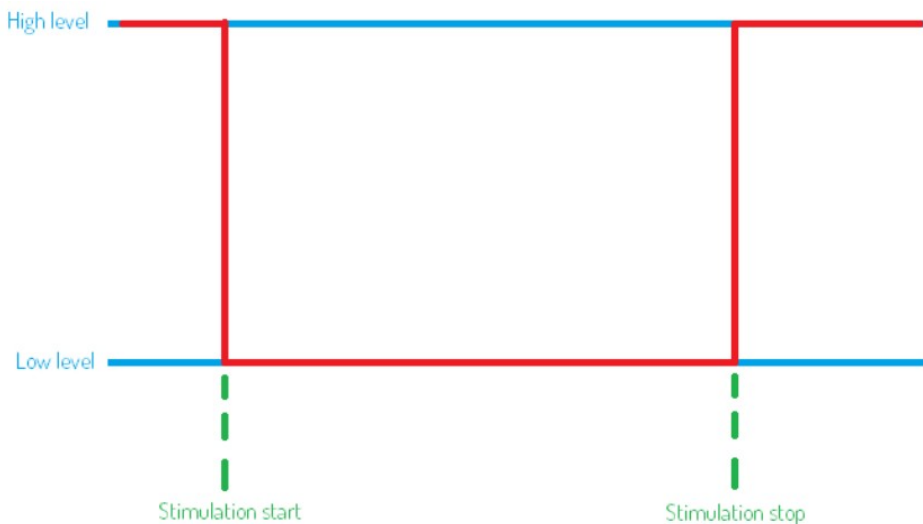


Figure 90. Trigger out setup for level low.

5.7.1 Available output events

- MAX (maximum) – A signal reached its maximum (only applicable to tACS) (Figure 91)
- MIN (minimum) – A signal reached its minimum (only applicable to tACS) (Figure 92)
- PHD – Phase detect, A signal reached given phase (only applicable to tACS) (Figure 93)
- ZERO (zero level) – A signal changed the polarity (only applicable to tACS) (Figure 94)
- NOC (number of cycles) – An event triggered following a specified number of signal cycles (only applicable to tACS) (Figure 95)
- PIT (point in time) – After a specified period of time, a trigger event occurs 'n' times with a predetermined interval between each other (Figure 96)
- STRT – Stimulation Started (Figure 97)
- STOP – Stimulation Stopped (Figure 97)

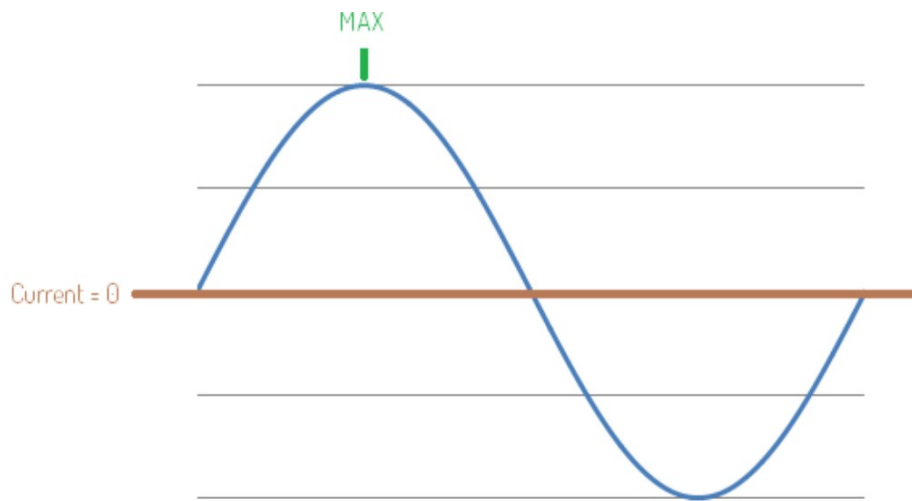


Figure 91. Trigger out event - signal maximum.

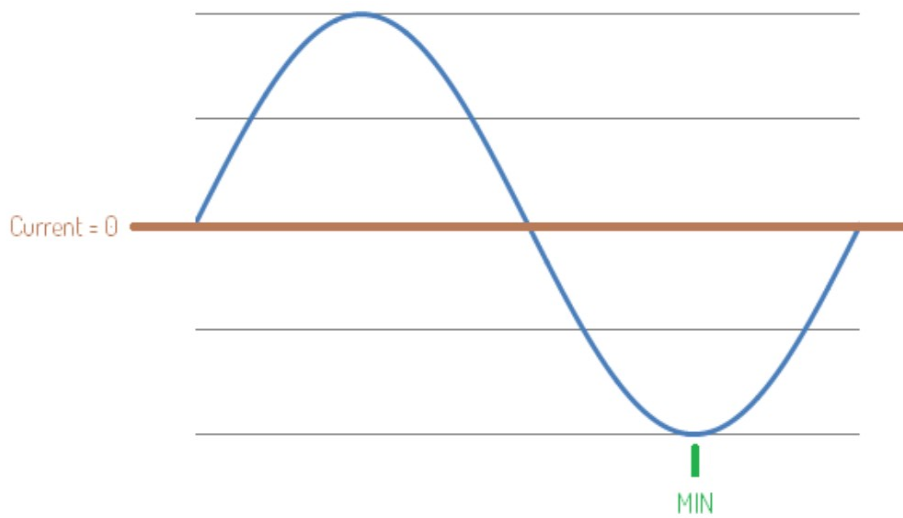


Figure 92. Trigger out event - signal minimum.

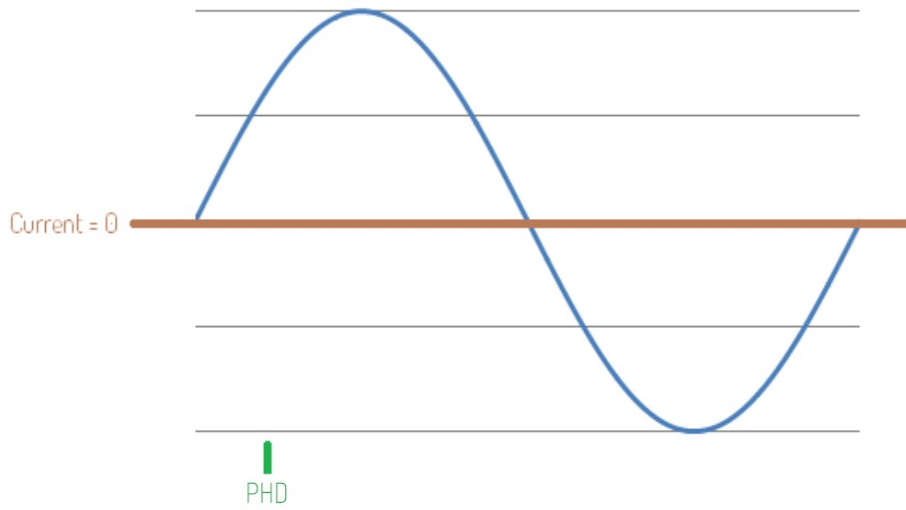


Figure 93. Phase detect trigger.

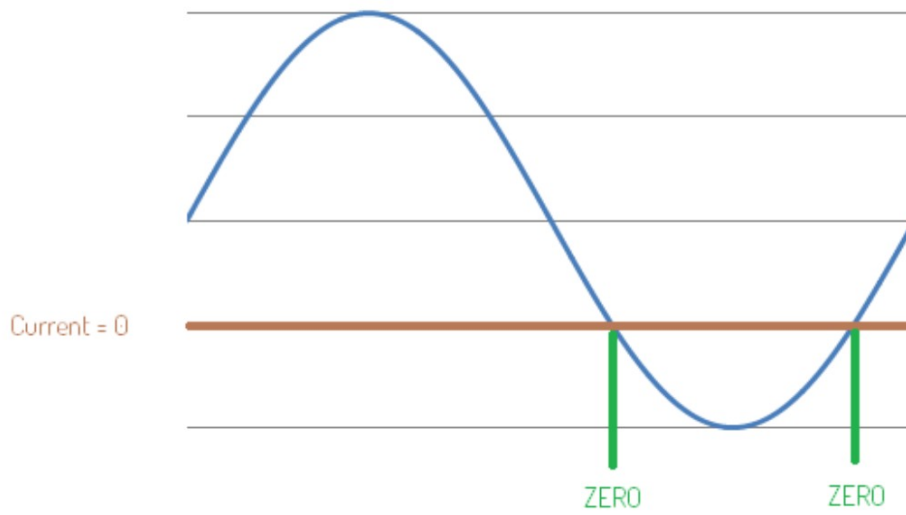


Figure 94. Trigger out event - zero level crossing

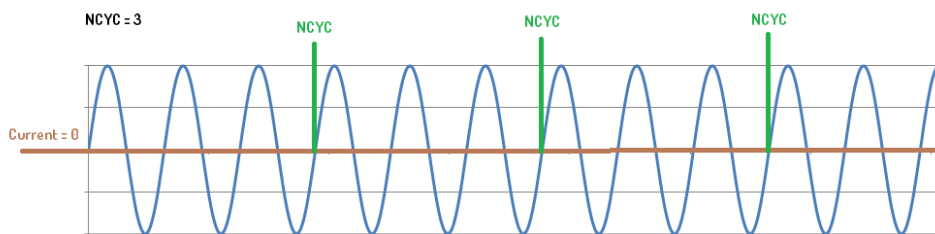


Figure 95. Trigger out event - number of cycles.

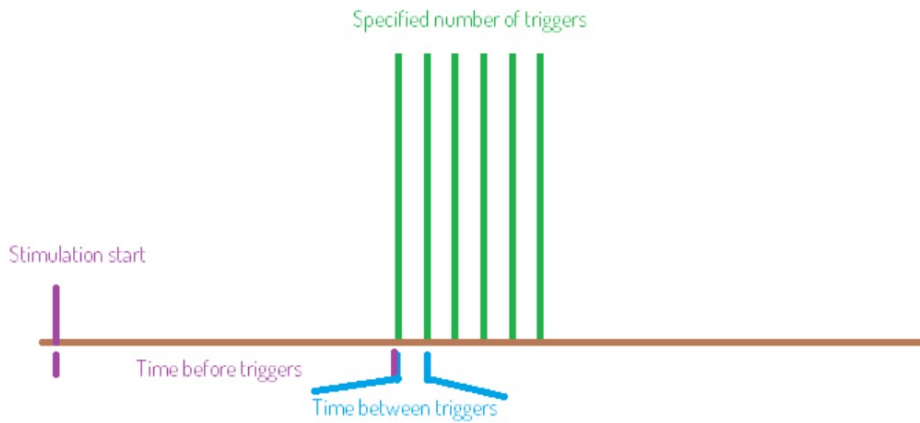


Figure 96. Trigger out event - point in time trigger.



Figure 97. Trigger out event - start and stop stimulation.

5.7.2 Triggers settings from device UI

To access the trigger settings, press the “Triggers” button on the stimulation parameter settings screen (see Figures 12-14). In this menu, the left column allows you to select the control signal to be configured. For input signals (trigger in), two channels are available: 1 and 2.

After selecting a channel, you can define the active signal edge (see Figures 84-86) by choosing the Rise and/or Fall, and then specify the stimulator’s response to the selected edge: Start and/or Stop stimulation.

All active interface elements (channel, edge, reactions) are highlighted in green. An example configuration is shown in Figure 98.

For output trigger settings, three channels are available. Each channel can be configured independently by selecting the output signal (see Figures 87-90). An example of the output trigger settings screen is shown in Figure 99. The first column of buttons in the configuration areas allows you to select the signal type. Only one signal type can be selected for each channel – the active option is highlighted in green. Depending on the stimulation mode and the selected signal type, different events may be available for generating an output signal. The widest range of events is available in tACS mode. Selected events are indicated by green highlighting of their labels.



Figure 98. Input triggers settings screen.

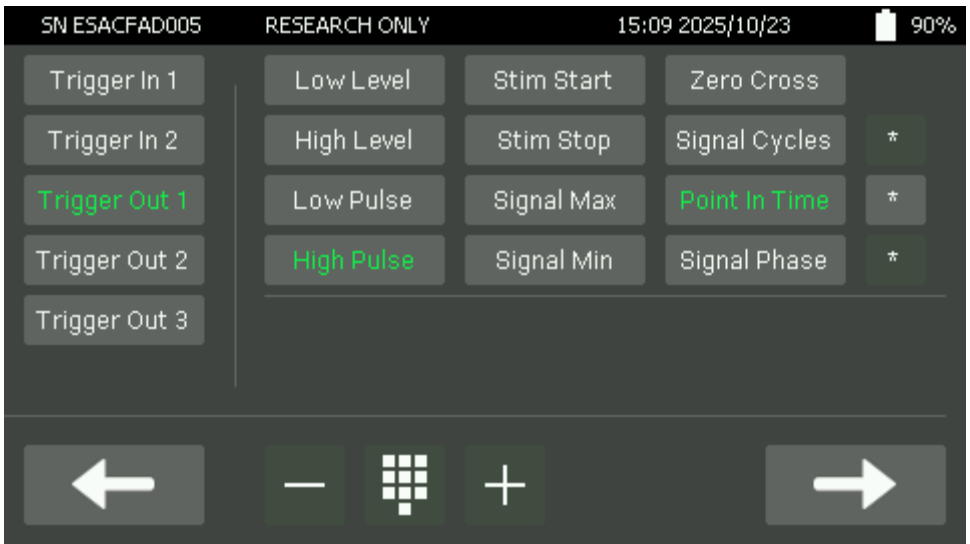


Figure 99. Output trigger settings screen for tACS mode. In tACS mode, all types of events that trigger the control signal are available for the Low/High Pulse output signal.

For the “Signal Cycles”, “Point in Time”, and “Signal Phase” events, additional parameters are available that define the behavior of the output signal. To access these parameters, activate the event and press the * button. The parameters that can be configured for the selected output trigger type will be displayed at the bottom of the screen. When you select the parameter, its label is highlighted in green, allowing you to enter a value using the numeric keypad, as described previously. An example screen showing additional parameters for the “Point in Time” trigger is shown in Figure 100.

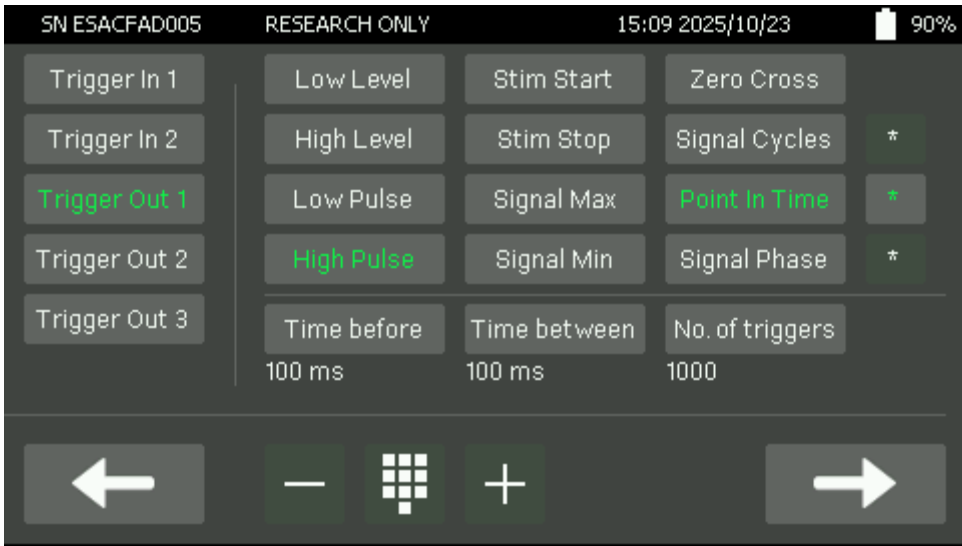


Figure 100. Output trigger settings screen with additional settings displayed for a Point In Time event.

Table 1 shows the permitted configurations of output triggers for individual stimulation modes and output signal types.

SIGNAL TYPE	EVENT	STIMULATION MODE			
		TDCS	TACS	TRNS	PULSE
LEVEL	START	✓	✓	✓	✓
	STOP	✓	✓	✓	✓
	MAX	✗	✗	✗	✗
	MIN	✗	✗	✗	✗
	ZERO	✗	✗	✗	✗
	NOC	✗	✗	✗	✗
	PIT	✗	✗	✗	✗
	PHD	✗	✗	✗	✗
PULSE	START	✓	✓	✓	✓
	STOP	✓	✓	✓	✓
	MAX	✗	✓	✗	✗
	MIN	✗	✓	✗	✗
	ZERO	✗	✓	✗	✗
	NOC	✗	✓	✗	✓
	PIT	✓	✓	✓	✓
	PHD	✗	✓	✗	✗

Table 1. Permitted trigger out signals depending on the type of stimulation and output signal.

6 Firmware update

The firmware of the device is constantly improved by a team of engineers. The device is analysed and user feedback is considered on an ongoing basis. To ensure a positive experience with the device users are offered free system updates.

6.1 Firmware distribution

Firmware update notifications will be sent to customers via email messages by the distributor. The message will contain a link to the file to be downloaded by the user.

The firmware update is distributed as a single IMG file.

6.2 Firmware update



WARNING! It is strongly recommended to use the dedicated application for firmware updates.

It is also possible to update the device using any serial terminal software that supports binary file transfer. This method requires familiarity with such software and the device's communication protocol. The firmware update process consists of two stages. The first stage involves uploading the file to the device using any serial terminal application. The second stage is flashing the device through the bootloader. While the second stage is performed automatically by the device, the first stage requires user interaction.

To upload new firmware to the device, connect it to a USB port. The device should appear as a new serial port. Connect to the device using a serial terminal that allows file transfers (NOTE: make sure to use a terminal that sends the file exactly as it is, without converting it to text or adding extra special characters). The connection parameters do not matter. To transfer the file, send the command: “:SYST:BOOT X”, followed by a carriage return, where X is the file size in bytes. If the device returns the message OK, it is ready to receive the file. The file can now be sent to the device. During the file upload, progress can be tracked on a bar displayed on the device's screen. Once the transfer is complete (the specified number of bytes has been sent), the device will reboot.

After the device restarts, firmware flashing begins. Upon successful update, a welcome screen will appear with the message “Update completed.”

The update process may take up to one minute. During the update, the device screen is turned off and the blue stimulation LED is on. WARNING: Do not turn off the device during the firmware update process, as this may render the device unusable.

6.3 Firmware Update Error Control

The device has strict firmware update error control. Several errors may occur during an update, due to hardware or software failures. Should this happen, an error message is displayed on the welcome screen.

List of error messages:

- ‘File with upgrade was corrupted. Upgrade aborted!’ – Uploaded file is incorrect or incomplete.
- ‘Uploaded firmware version is older or does not match hardware. Upgrade aborted!’ – Wrong version of firmware was uploaded. It is not applicable to the user's hardware or is older than the current firmware.
- ‘Error occurred while updating. Upgrade aborted!’ – Internal error. If it is recurring, contact the distributor – the device is malfunctioning.
- ‘Firmware in device was corrupted. Backup restored.’ – The device discovered corruption in its firmware and restored a backup program – WARNING: The backup program is the previous working version of firmware!

In case of an error, the previous firmware version is restored.

If the error is due to malfunction the device has no means to start a correct program, the blue Stimulation LED starts to blink at device start-up. When this happens, contact your distributor.

7 Troubleshooting and maintenance

7.1 Device-specific issues

7.1.1 Device interface

1. The device screen is not working
 - The device is shut down automatically for the duration of charging
2. The device screen is not responding
 - The operator is wearing gloves.

7.1.2 Stimulation

1. Stimulation does not start exactly on pressing the 'START' button.
 - There is a delay before proper stimulation starts. An impedance measurement is optionally performed before starting the stimulation, followed by a momentary delay before the diagnostic procedure begins.
2. Stimulation does not start (wrong impedance message dialog is displayed).
 - Verify the positioning of the electrodes, check electrode connections, check or remove adapters if any were used to connect the electrodes.
3. User-defined impedance cannot be set to a higher value.
 - User-defined impedance must always be lower than the security threshold, which can also be set in a limited range (not exceeding the maximum of 50k Ω).
4. The device stops working during stimulation.
 - Electrode connection conditions have changed, impedance has exceeded the acceptable value, e.g. exceeded 50k Ω or dropped below 300 Ω .
5. 'Parameters unreachable' message is displayed.
 - This message indicates that the impedance is so high that the current settings cannot be achieved due to hardware and safety limitations.
6. A sound signal is emitted
 - This is normal activity of the device, indicating that the user-defined impedance level has been exceeded or another stimulation error occurred (check the messages on the device screen or the status received using SCPI protocol).

7.1.3 Device operation

1. The charging LED remains on, despite that 15 hours have passed after connecting the device to a charger.
 - Disconnect the device from the charger immediately and contact your distributor.
2. Battery charging indicator LED is continuously on.
 - Do not connect the device to a charger, contact your distributor.
3. Battery empty (5%).
 - Connect the device to a charger. (connecting USB does NOT result in charging!)

4. Charging indicator LED is off.

- Check the charger connection, use an original charger, the device will not charge when connected to a PC via SCPI protocol.

7.2 Maintenance

The device is designed for self-checking its critical functions and will alert if manufacturer's service is required.

If a critical function of the device is non-functional, the device will lock and alert on service needed.

There's no need for a scheduled maintenance or re-calibration.

Any service actions within the device should be performed by the manufacturer's service and engineering department or the authorized and trained representative of the manufacturer only.

There are no user-serviceable parts in the device.

All safety fuses inside the device are fixed to internal printed circuit boards and are not replaceable or resettable.

Internal batteries are in-built and are not replaceable by the user. In case of an evident loss of operational time due to battery aging, the device should be serviced by the authorized service personnel only.

Any wear and tear of the accessories, such as cables and electrodes should be noted and a new accessory should be ordered to replace a worn one.

Electrodes, charger and some cables are off-the-shelf parts available on the market and may be replaced by a new one of exactly the same model by the same manufacturer.

All worn or damaged cables must be re-ordered from the manufacturer.

Using any replacement parts other than those specified in this manual is prohibited.

8 Manufacturer and servicing information

The MANUFACTURER will make available on request: circuit diagrams, component part lists, descriptions, calibration instructions, or other information that will assist SERVICE PERSONNEL to repair those parts of ME EQUIPMENT that are designated by the MANUFACTURER as repairable by SERVICE PERSONNEL.

SERVICE PERSONNEL must be an authorized and trained representative of the manufacturer or the manufacturer's service or engineering department.

The manufacturer and support provider for the medical device is:

QVITI S.A.

Address: Warszawa 01-552, pl. Inwalidów 10

Telephone: + 48 690 315 745

kontakt@qviti.com

<https://www.qviti.com/>

9 Revision history

Revision	Date	Authors	Description
1.0	18.02.2025	MO, SM, SS.	First release
1.1	20.10.2025	MO	Manual update for compatibility with firmware version x.1.0 and company name change.