mTrigger® Biofeedback System User Manual

Please read the entire User Manual before attempting to operate this device.

If you have any questions or problems with this device, please contact:

mTrigger® Customer Service:
550 South College Avenue
Suite 110
Newark, DE 19713

US and Canada: *
Email: info@mtrigger.com

Manual Part Numbers:
863712000337 - Individual Unit + Accessories
863712000382 - Clinical Bundle

Effective Date: 1 November 2017
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</table>
1 | Glossary andAbbreviations

Table 1.1: Glossary

Table 1.2: Abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Term</th>
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</thead>
<tbody>
<tr>
<td>EMC</td>
<td>Electromagnetic Compatibility</td>
</tr>
<tr>
<td>FDA</td>
<td>Food and Drug Administration</td>
</tr>
<tr>
<td>US or USA</td>
<td>United States</td>
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</table>

2 | Safety
This section provides safety guidelines and safety-related statements to operate this biofeedback system safely and effectively. Additional guidelines, statements, and protocols appear throughout this manual. Follow all printed guidelines, warnings, cautionary statements, and protocols when using this biofeedback system.

2.1: Conventions Used in this Manual
Various Warnings, Cautions, Recommendations and Notes are presented throughout this manual. Explanations and the corresponding symbols are:

**Warning**: Specific or potential danger. If ignored or compromised, the situation could result in harm. Warning statements are preceded with a yellow symbol.

**Caution**: Possible problem with the device associated with its use or misuse. Problems include, but are not limited to, device malfunction, device failure, or damage to the device. Caution statements are preceded by a black-and-white symbol.

**Recommendation**: Offers guidance for the optimal application and usage of the device. Recommendation statements are in a shaded box.

**Note**: Describes the conditions or exceptions that may apply to the subject matter presented.
3 | System Set-Up

3.1: Location
The mTrigger\textsuperscript{®} Biofeedback System can be used in virtually any location, so long as there is sufficient space to perform appropriate therapy exercises. Locations with high levels of electrical activity may cause minor signal interference, but do not pose risks to the system or users.

3.2: Unpacking and Initial Set-up

Step 1. Using the mTrigger\textsuperscript{®} Biofeedback System packing list, carefully unpack the system and its accessories from the shipping box.

Step 2. Check for any missing or damaged parts.

Step 3. If items are missing or damaged, call mTrigger\textsuperscript{®} Customer Service. Notify the carrier if the damage occurred during shipping and retain shipping packaging for evidence.

Step 4. Fully charge the battery before using the device(s) via the USB port for the first charge. See figure 3.1

Fig. 3.1: Micro USB charging port setup

Fig 3.3: Charging state LEDs
green = fully charged, red = actively charging, white = not charging/in use
**Recommendation:** Charge the device before use. The system ships with the device battery only partially charged.

**Recommendation:** Turn devices off before any method of charging. When the device is on, the LEDs cannot indicate a full charge and full charge will take longer.

### 3.3: Battery Use and Charging

**Warning:** Do not modify the battery or attempt to remove it from device.

**Warning:** Do not immerse the battery in water or get the battery wet.

**Caution:** DO NOT leave device charging overnight; charging time in excess of 6 hours can cause over-charging and impact battery life and device performance.

**Caution:** The temperature range for charging this battery is -10° C to 45° C (14° F to 113° F). Do not charge, store, or attempt to use the battery outside of this range.

**Caution:** Do not store the battery over a long period of time at temperatures above 30° C (86 °F), such as inside a car on a hot day or in direct sunlight; this may damage the battery.

**Caution:** Do not disassemble or open, crush, bend, deform, puncture, shred or burn the battery.
**Note:** When the device is plugged in or wirelessly charging, it will be illuminated in red. Green illumination indicates full charge. When in use, it will be illuminated in white. See figure 3.3.

### 3.4: Battery Malfunction

If the battery is not charging as expected, contact mTrigger® Customer Service to determine if a replacement unit is needed. The battery cannot be removed from the unit or replaced except by the manufacturer.
4 | Device Description

The mTrigger® biofeedback system includes:

- *mTrigger* biofeedback device (qty 1 for Individual Unit; qty 3 for Clinical Bundle)
- Sensing cables (qty 2, IU; qty 6, CB)
- Sensing electrodes (qty 10 pouches, IU; qty 50 pouches, CB)
- User interface software application (free download to mobile device)

A custom user interface software application allows the operator to pair with a specific device via Bluetooth and set exercise parameters, complete training sessions, track progress, play games, and calculate neuromuscular deficit. Once the operator has selected the desired exercise parameters, complete training session and save output to the tracking module. These steps are detailed in Section 5.

**Warning:** Do not use any accessories and/or cables not specified or sold by mTrigger® as replacement parts. Use of unauthorized accessories and/or electrodes not specified or sold for use with this system may result in ineffective signal transmission or damage to the system.

**Warning:** DO NOT attempt to gain access to any internal component. Doing so may cause injury and/or device damage. THERE ARE NO USER-SERVICEABLE COMPONENTS inside this device.

4.1: Indications for Use (Purpose of the Device)

This biofeedback system is indicated for:

- Biofeedback, muscle re-education
- Deterrence of atrophy due to lack of voluntary muscle activation
- Increasing voluntary muscle control and activation via real time display of muscle activity
- Stroke rehab by muscle re-education
- Visual feedback to supplement physical therapy exercises performed for the purpose of rehabilitating musculoskeletal injury
4.2: Contraindications

None.

Caution: Use of the mTrigger® Biofeedback System simultaneously with an electrical muscle stimulation system may impact signal quality.

4.3: Biofeedback System Unit

Top View: 

Bottom View: 

Right Side View:

Power Switch (position: off)

Serial Number
Left Side View:

<table>
<thead>
<tr>
<th>1</th>
<th>2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Channel 1 EMG port</td>
<td>Channel 2 EMG port</td>
</tr>
<tr>
<td>Micro USB charging port</td>
<td></td>
</tr>
</tbody>
</table>

4.4: Accessories

**Warning**: Use of unauthorized accessories and/or electrodes not specified or sold for use with this system may result in ineffective signal transmission or damage to the system.

**Micro USB Charger:**

![Micro USB Charger](image)

**AC Power Cord:**

![AC Power Cord](image)
Sensing Electrodes:

Sensing Cables:

Electrode connector pins

EMG port jack

Cable connector receptor
5 | Operation

5.1: Safety Precautions

Caution: FAILURE TO COMPLY with the setup procedures and suggested application techniques listed in the manual may lead to ineffective treatment via improper signal transmission or device failure.

Take the following steps to secure the treatment area:

- Clear exercise area of any obstacles.
- If exercise is stationary, place device on a nearby table or other flat surface where it will not impede treatment. Prop phone or tablet on provided viewing stand where the application can be easily seen by the patient and provider, if desired.
- If exercise is mobile, secure the device in the mTrigger wearable band or a pocket; if neither option is available, hold the unit during exercise.

Note: For mobile exercises, either prop phone or tablet on provided viewing stand where the application can be easily seen by the patient and provider, or hold mobile device in hand.

Caution: DO NOT rest mTrigger unit on or touching other electronic devices (computers, printers, digital scales, etc.) as this will cause electronic interference and improper signal transmission.

5.2: Patient Preparation

Warning: DO NOT use in conjunction with electrical stimulation treatment. This could cause device damage or malfunction.

Before a treatment, perform the following preparation:

- Ensure that the skin on the treatment area is clean, dry, and free of surface dirt or oils for proper electrode adherence and signal reception. See instructions on electrode packaging.

5.3: Device Operation

Additional content: Quick Startup Guide (included in package or download from website)

1) Search “mTrigger biofeedback” on the Apple Store and download the mTrigger Biofeedback App to your mobile device. App is available across Android and iPhone phones and tablets.
2) Attach electrode connector cable(s) to mTrigger EMG ports.

![Attaching electrode connector](image)

**Caution:** For single channel use, only data from Channel 1 will be read. Please insert channel 2 cable ONLY for dual channel use. Inserting Channel 2 cable during single channel use could impact signal quality.

3) Attach sensing electrodes to sensing cables

![Attaching sensing electrodes](image)

4) Place one pair of electrodes on active muscle. See Electrode Placement Database for suggested electrode placements.

**Note:** For dual channel use, i.e. monitoring of two muscular activity channels, place second electrode pair on a co-contracting muscle or corresponding muscle on the healthy side.

5) Turn on device by sliding power switch on the right hand side as indicated. A white light will illuminate the m when the device is on.
Note: If charging is necessary, any micro USB charger will work; active charging is indicated by a red light, charge complete is indicated by a green light. Very low battery level may impact the unit’s ability to communicate data over the BlueTooth connection.

6) Open application on mobile device.

7) Select “CONNECT” in the upper right corner of the Home screen, or access “ADJUST BLUETOOTH SETTINGS” on the Settings page.
8) Locate the serial number on the sticker on the underside of the mTrigger device. Match the serial number to the unit numbers that appear in the scanned device list. Tap to select your device number and connect via BlueTooth. A “device connected” confirmation will pop up.

9) From the Home screen, go to Settings to set up treatment parameters:
a) Channel Select: Select dual or single channel as desired.
   i) If in single channel mode, ensure you are plugged into Channel 1 only; Channel 2 will not display data in single channel mode, but unintended Channel 2 inputs could impact the Channel 1 signal quality.

b) Time Settings: Modify total, flex, and relax time as desired. Both flex and relax times must be set or you will be notified of an error. Reps are calculated as \([\text{total time}] / [\text{flex time} + \text{relax time}]\). Every training session will begin with the relax period of the rep.

c) Channel Goal Settings: Set the microvolt goal for each channel; this can also be set in Train by testing in real time to assess MVC. Goal can also be modified by typing the exact desired uV level into the text bar (tap on the number before uV to type).

**Note:** 1500uV is usually a good starting point to assess patient capability

10) Save settings OR tap “BACK” in upper left or \(m\) in lower left to return to home screen and accept “save settings” prompt upon exiting
11) From the Home screen, tap to enter Train.

a) Before beginning your session, modify the goal as necessary to match MVC using the slider bars or by typing.

b) Hit the play button on the bottom bar to begin your session.

c) Pause button pauses the session and will restart at the beginning of the relax time of the previous rep.
d) Save results when prompted at session completion to see timestamped output in the Track module.

12) To Play: Tap Play from Home to launch the game list.

a) Select the VIEW [GAME NAME] button to take you to the individual app for that game. There you can customize settings for that session of play; universal app settings will not be affected.
13) Neuromuscular Deficit Test: First, enter settings and toggle to dual channel mode. Then ensure that the goals for CH1 and CH2 match exactly – you can tap the numbers to type. The goal should be set to both at the level of a normal max contraction on the healthy side to provide the most accurate deficit calculation.

Tap Neuromuscular Deficit Test to start the test. NDT assesses the deficit of the involved side as compared to the healthy side. The involved side will test on the left side of the screen (CH1), normal side on the right (CH2). Flex your involved muscle when indicated by the green “I” and left-pointing arrow; flex your normal muscle when indicated by the green “N” and right-pointing arrow. The test compares average microvolt activity on the healthy versus involved sides over a one-minute test period. See help button on bottom right of screen for further details.

Note: Ensure that your involved muscle is plugged into channel 1 during the NMDT. It is also critical that electrode placement be as identical as possible on both healthy and involved sides of the body. Ensure that goals are set exactly equal to one another in settings. Failure of any of these steps will invalidate test results.

14) Tap Track and select a session to see actual EMG activity compared to goal per timestamped session. Tap an individual graph to see average maximum contraction for that session.
15) Slide the power button up to shut down the device. Bluetooth will disconnect automatically upon powering device off or force closing app.

6| Maintenance and Calibration

**Warning:** DO NOT attempt to gain access to any internal component. Doing so may cause injury and/or device damage. THERE ARE NO USER-SERVICEABLE COMPONENTS inside this device.

6.1: Cleaning

**Warning:** Always turn off the device before cleaning and remove from charger.

Keep EMG ports and sensing cable jacks clear from dirt, dust, and debris. These substances will impact signal quality and device functionality and over the long term could cause device damage.

If cleaning is necessary, use a dusting cloth.

**Warning:** DO NOT use water or a damp cloth to clean any part of the mTrigger Biofeedback System; no component of the system is waterproof.
Caution: Water coming in contact with sensing electrodes may impact skin conductance and thereby signal quality.

6.2: Before You Call—Troubleshooting

If you are having a problem with your biofeedback system, please check the list below of common conditions that can occur that you may be able to resolve without having to contact mTrigger® Customer Service.

If the device will not turn on:
Attempt to charge. If after attempted charge, the device does not show charging indicator LEDs or still does not turn on, call mTrigger® Customer Service to undergo troubleshooting exercises to determine if a replacement battery is required.

If there are problems with the battery:
Problems with the battery (e.g., battery will not charge, will not hold a charge, or depletes charge quickly) may not require battery replacement. Call mTrigger® Customer Service to undergo troubleshooting exercises to determine if a replacement battery is required.

If there are error messages:
Ensure that your phone’s Bluetooth is turned on and that you are using only authorized accessories.

7 | Quality and Regulatory

7.1: Quality Environment
- The mTrigger® biofeedback system production environment complies with 21CFR820.70
- The mTrigger® biofeedback system is manufactured in compliance with the United States Food and drug Administration’s general manufacturing practices concerning medical devices.

7.2: Regulatory Compliance
According to the applicable standards, the mTrigger™ Biofeedback System is classified as follows:
- Class II Medical Device per USFDA 21CFR 882.5050
- 510(k) exempt
In order to safely perform its intended use, the system requires two accessories whose classifications and registrations are as follows:
• Custom Sensing Electrodes – Cutaneous electrodes classified as Class II (special controls; 510(k) exempt) per USFDA 21CFR Sec. 882.1320
• Custom Sensing Cables – not classified as a medical device or medical device accessory; tested as part of the mTrigger® Biofeedback System EMC and safety standards

Please see supplemental FDA Compliance Summary Report (Quality Document number Q-06) for further details.

8 | Specifications and Environmental Conditions

8.1: Specifications

Table 8.1 System Specifications

<table>
<thead>
<tr>
<th>Model Name</th>
<th>mTrigger® Biofeedback System</th>
</tr>
</thead>
<tbody>
<tr>
<td>Model Number(s)</td>
<td>863712000337 - Individual Unit + Accessories</td>
</tr>
<tr>
<td></td>
<td>863712000382 - Clinical Bundle</td>
</tr>
<tr>
<td>Device Classifications</td>
<td></td>
</tr>
<tr>
<td>United States</td>
<td>Class II Medical Device per USFDA 21CFR 882.5050</td>
</tr>
<tr>
<td>User Interface</td>
<td></td>
</tr>
<tr>
<td>Mobile Application</td>
<td>mTrigger® Biofeedback</td>
</tr>
<tr>
<td>Mechanical Specifications</td>
<td></td>
</tr>
<tr>
<td>Dimensions</td>
<td>4 in x 2.4 in x 1.1 in</td>
</tr>
<tr>
<td>Weight</td>
<td>0.25 lb</td>
</tr>
<tr>
<td>Environmental Specifications</td>
<td></td>
</tr>
<tr>
<td>Operation temperature</td>
<td>10 °C to 35 °C │ 50 °F to 95 °F</td>
</tr>
<tr>
<td>Storage temperature</td>
<td>-20 °C to 70 °C │ -4 °F to 158 °F</td>
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<tr>
<td>Electrical Specifications</td>
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<tr>
<td>Input Voltage</td>
<td>TBD in EMC testing</td>
</tr>
<tr>
<td>Input Current</td>
<td>TBD in EMC testing</td>
</tr>
</tbody>
</table>
8.2: Accessory Details

Warnings & Precautions: mTrigger Custom Sensing Electrodes

- Keep electrodes out of reach of children.
- Electrodes are intended for single person use only.
- DO NOT remove electrode by pulling on the lead wire.
- DO NOT use generic electrodes with the mTrigger Biofeedback System, as damage to mTrigger accessories will occur.
- DO NOT use electrodes that have been incorrectly applied or damaged.
- Follow placement instructions carefully – electrodes should be separated on the skin and positioned as indicated for your treatment.
- DO NOT apply to broken skin. Should a skin rash or irritation occur, discontinue use and contact your physician.
- DO NOT use the mTrigger Biofeedback System while operating machinery, sleeping, or in close proximity to water.

Note: The life of electrode adhesive gel varies depending on skin condition, preparation, storage, and climate. Please refer to care and storage instructions on the electrode packaging.

Table 8.2 Accessory Specifications

<table>
<thead>
<tr>
<th>Device Classifications</th>
<th>Item</th>
<th>Description</th>
<th>Classification</th>
<th>Image</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Sensing Electrodes</td>
<td>2&quot; square white cloth, sensing gel, self-adhering surface electrodes</td>
<td>Class II special controls; 510(k) exempt</td>
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</tr>
<tr>
<td></td>
<td>Sensing Cables</td>
<td>4’ M/F AUX jack pin connector pigtail cable</td>
<td>Non-medical</td>
<td></td>
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</tbody>
</table>
9 | Warranty Information

9.1: Terms and Conditions
The mTrigger Biofeedback system is warranted to be free from manufacturer defects in materials and workmanship for a period of 90 days, starting from the date of initial shipment. This warranty does not extend to incidental or consequential damages nor to damage caused by negligent or improper handling in use or storage; products on which the original identification markings or labels have been intentionally defaced, altered or removed are not eligible for warranty.

mTrigger, LLC reserves the rights of determining existence and cause of defect as well as the option to repair or replace products which prove to be defective as necessary during the warranty period. Claims may be submitted for review after the warranty period close; however, mTrigger does not guarantee repair or replacement after the close of the warranty period.
This warranty extends only to the original purchaser of the equipment from mTrigger, LLC. The purchaser must notify mTrigger within 15 days of first detecting the defect and promptly return the defective product before expiration of the warranty period.

Return shipping costs will be covered by mTrigger, LLC within the warranty period. mTrigger will return repaired or replaced products to purchaser within the Continental United States at no additional charge. International transportation fees and insurance (if necessary) beyond this limit will be charged to purchaser.

9.2: Return Procedure
Step 1. Review the date of shipment to determine the validity of any warranty claim. Warranty claims should only be made for products within the terms of the warranty policy (see Section 9.1).

Step 2. Before contacting mTrigger Customer Service, be prepared to provide:
• Product Model and Serial Number (see Section 11.1)
• Purchase and Shipment Date
• Invoice number
• Reason for Return
• Contact name, phone number, and email address for further communication

Note: Providing complete information as requested will expedite the procedure.

Step 3. Call or email Customer Service to obtain a Return Material Authorization (RMA) number and detailed return instructions. Customer Service will email a form to you.

Step 4. Complete and sign the RMA form and return to mTrigger Customer Service. Adhere to mTrigger’s complete return instructions for transportation and packaging
to ship the product (freight and insurance prepaid) with completed and signed RMA form to mTrigger.

**Step 6.** Once MTrigger receives and evaluates the product, MTrigger will advise the purchaser of the warranty service determination and any subsequent action, i.e. shipment of repair or replacement unit(s).

10 | **Labeling**

The label displays the model number and serial number, which encodes the date of manufacture as the lot number as well as the individual device’s Bluetooth pairing number.

<table>
<thead>
<tr>
<th>Position</th>
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<th>Alpha/Num</th>
<th>Notes</th>
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</thead>
<tbody>
<tr>
<td>1</td>
<td>product key</td>
<td>1</td>
<td>alpha</td>
<td>M = main unit</td>
</tr>
<tr>
<td>2</td>
<td>Rev #</td>
<td>2</td>
<td>num</td>
<td>no punctuation</td>
</tr>
<tr>
<td>3</td>
<td>manufacturing date/lot #</td>
<td>4</td>
<td>num</td>
<td>yy/ww</td>
</tr>
<tr>
<td>4</td>
<td>board serial #</td>
<td>3</td>
<td>num</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Product Key</th>
<th>PCB rev #, no punctuation</th>
<th>Production date yy/ww</th>
<th>PCB serial #</th>
</tr>
</thead>
</table>
11 Contact Information

11.1 Customer Service
For immediate assistance, contact mTrigger® Customer Service directly.

If this biofeedback system does not function as expected and/or if the biofeedback system malfunctions, contact mTrigger Customer Service.

Please have the device serial number so that Customer Service can provide you with the highest level of service.

The Serial Number (SN) of the device are found on the black barcode sticker located on the bottom of the unit.

mTrigger Customer Service:
550 S College Ave
Suite 110
Newark, DE 19713

Phone: 1-877-627-3858 ext. 1
Email: info@mtrigger.com

Sales & Other inquiries:
Phone: 1-302-502-7262 ext. 0
Email: info@mtrigger.com