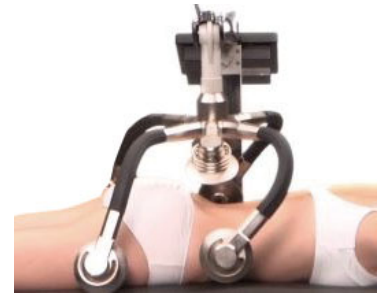




**ERCHONIA**  
World Leaders in Low Level Laser Technology



## Non-Prescriptive/Non-invasive Fat Loss Laser



## INSTALLATION AND PROPER USE REFERENCE GUIDE

**Read this entire booklet before using your ZERONA Z6 OTC Laser**



# READ THIS FIRST

To ensure proper use, and to achieve your best results, it is important that you read and understand the instructions, warnings, precautions and safety information in this booklet before using your Zerona Z6 OTC laser for the first time.



This symbol appears next to information about possible safety risks.

Questions? Our Erchonia Customer Care representatives are available to help. Contact us at:

**Erchonia Customer Care**

Phone: 1-321-473-1251

Email: [info@erchonia.com](mailto:info@erchonia.com)

Or visit [erchonia.com](http://erchonia.com)

**Erchonia Europe Customer Care**

Phone: +44 (0)1491 821135

Email: [info@erchoniaeurope.com](mailto:info@erchoniaeurope.com)

Or visit [erchonia.com](http://erchonia.com)

**ATTENTION:** This device is sold “Over the Counter” (OTC). In general, there is no prescription required for the use of this device by any individual. Notwithstanding the foregoing, if you are a licensed healthcare professional using this device, you are solely responsible for your use of this device and for ensuring that this device is properly used in accordance with any professional or other regulations that may apply to your use of this device. The device should always be used in accordance with the instructions set forth in this manual.

We recommend you periodically contact Erchonia Corporation to determine if additional product information updates are available.

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# SECTION 1: GENERAL INFORMATION

## HOW TO USE THIS REFERENCE GUIDE

This reference guide is designed to take you through set-up and proper use of your Zerona Z6 OTC laser. It will cover Warnings and Precautions, Assembly, Proper Placement of Laser Diodes, Treatment Protocol and Client Measurement Guidelines.

It is important that you read and understand all the information contained in this reference guide before doing any treatments with the Zerona Z6 OTC laser. If you have any questions, contact our Erchonia Customer Care representatives.

## WHAT IS THE ZERONA Z6 OTC LASER?

The ZERONA Z6 OTC Laser is indicated for use as a non-invasive dermatological aesthetic treatment for the reduction of body circumference.

The Zerona Z6 OTC laser delivers a new non-invasive fat loss procedure designed to slim the body without surgery, pain or needles. The protocol allows clients to continue daily activity without interruption.

Clinical data on safety and effectiveness has only been generated for waist, hips, thighs, upper abdomen, and upper arms. Presently there does not exist clinical data on safety or effectiveness for other parts of the body

## HOW DOES IT WORK?

The Zerona Z6 OTC laser works by using a patented and clinically proven low-level laser technology. Each laser diode of the device emits visible red light (about 635 nanometer wavelength). The laser emulsifies the fat, (defined as leakage of the fat out of the cell) within the adipose tissue which is then released into the interstitial space. The excess fat is then passed through the body during its normal course of detoxification.

# PACKAGE CONTENTS














The Zerona Z6 OTC laser is shipped in one large freight box with three components:

- Z6 control base
- Z6 laser diode arm
- Accessories (small box)
  - Safety glasses
  - Tape measure
  - Keys
  - Manual

When you receive the shipment, carefully inspect the container for damage. If the shipping container or cushion material is damaged, keep it until the contents have been checked for completeness and the device has been checked for proper function. If the contents are incomplete or if there is mechanical damage, contact Erchonia Corporation at 1-321-473-1251. If the shipping container is damaged, also notify the carrier.

# SYMBOLS USED ON THE EQUIPMENT

Any or all of the following symbols may be used in this reference guide or on the equipment:

Symbol	Description
	Temperature Limitation
	Type B receipt connection - applied parts that are generally not conductive and can be immediately released from the client.
	Conformité Européenne - Complies with EMC 2004/108/EC and LVD 2006/95/EC.
	Power ON/OFF
	Date of Manufacture
	Manufacturer
	Authorized representative in the European Community.
	Refer to Operating Instructions.
	<b>Warning</b> alerts you about a situation which, if not avoided, could result in death or serious injury. It may also describe potential serious adverse reactions and safety hazards.
	<b>Precaution</b> is used for the statement of a hazard alert that warns you of a potentially hazardous situation which, if not avoided, may result in minor or moderate injury to the user or client or damage to the equipment or other property. It may also be used to alert against unsafe practices. This includes the special care necessary for the safe and effective use of the device and the care necessary to avoid damage to a device that may occur as a result of use or misuse.
	No pushing.
	Pinch point-keep hands clear
	WEEE Symbol- Separate Collection Recommended

# SAFETY INFORMATION

Read the following important safety information before using the Zerona Z6 OTC laser.

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## WHEN NOT TO USE

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Avoid using the laser over a pregnant uterus.



DO NOT treat the face, neck or breast.

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## WARNINGS

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You must follow these Instructions for Use when using the Zerona Z6 OTC laser. Not following these instructions may result in serious injury.



**DO NOT** place/operate this device near mirrors or other reflective materials or surfaces as the laser light could be deflected off a mirror or reflective surface into someone's eyes, possibly causing eye damage.



**DO NOT** disassemble, modify, or remodel the device or accessories. This may cause device damage, malfunction, electrical shock, fire, or personal injury.



To avoid risk of electric shock, do not defeat the safety purpose of the grounding-type plug. A grounding-type plug has two blades and a third grounding prong. The wide blade or third prong is provided for your safety. If the provided plug does not fit into your outlet, consult an electrician.



Avoid direct contact of the laser beam into the eyes. Laser protective eyewear should be worn by the client to block light energy from the eyes during treatment.



**Beware of electrocution.** Do not submerge any part of the device in water. This could damage the device, or cause an electric shock that may lead to serious injury or death.



Use of controls or adjustments or performance of procedures other than those specified in this booklet may result in hazardous radiation exposure.



The Zerona Z6 OTC laser is not intended for use by persons with reduced physical, sensory or mental capabilities.

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## WARNINGS (Continued)

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Keep out of reach from children



Dispose of device in accordance with local and national regulations and codes. When spent and beyond repair or functional use, the device can be sent back to the manufacture for disposal. This ensures the proper separation and handling of all the internal parts and reduces any risk to the end user and the environment.



**DO NOT** permit any foreign materials or liquids to enter the device. Take care to prevent any foreign materials including, but not limited to, inflammables, water, and metallic objects from entering the device. These may cause device damage, malfunction, electrical shock, fire, or personal injury. To achieve the specified level of protection against spilled or splashed liquids, unplug the device from the power supply and thoroughly dry all exposed surfaces of this device and allow to dry thoroughly prior to operation.



Avoid using the laser on skin that is infected, burned or cut, as this could cause discomfort or irritation. Wait until the infection, burn or cut has healed before using the laser. If the client has a skin condition, lack of skin sensitivity or are undergoing professional cosmetic treatments, please check with their doctor or clinician before using the laser.”



Laser treatment should not be applied over, or in proximity to (near), cancerous lesions (such as moles and other suspicious skin markings unless known not to be cancerous). There have not been conclusive tests that show if low level laser exposures on or near cancerous lesions do or do not create a serious adverse effect.

## PRECAUTIONS



Read, understand, and practice the precautionary and operating instructions. Know the limitations and hazards associated with using any laser device. Observe the precautionary and operational decals placed on the device. If you have difficulty operating the device after carefully reviewing this reference guide, contact our Erchonia Customer Care representatives for assistance.



**DO NOT** position equipment so that in an emergency it is difficult to disconnect the power cord from the electric supply.



**DO NOT** use sharp objects such as a pencil point or ballpoint pen to operate the buttons on the touch screen as damage may result.



**DO NOT** place/operate this device in close proximity (6 inches) to other devices that emit frequency. If this device causes interference to other devices, which can be determined by turning the device off and on, you are encouraged to try to correct the interference by one of the following measures:

- Reorient or relocate the receiving device.
- Increase the separation between the equipment
- Connect the equipment into an outlet on a circuit different from that to which the other device(s) are connected.



This device should be operated in temperatures between 59° to 85° F (15° to 29° C), and transported in temperatures between -22° to 158° F (-30° to 70° C), with relative humidity less than 75%.



There are no user-serviceable parts inside the device except for the fuses. If a malfunction occurs, discontinue use immediately and contact Erchonia Corporation for repair service.



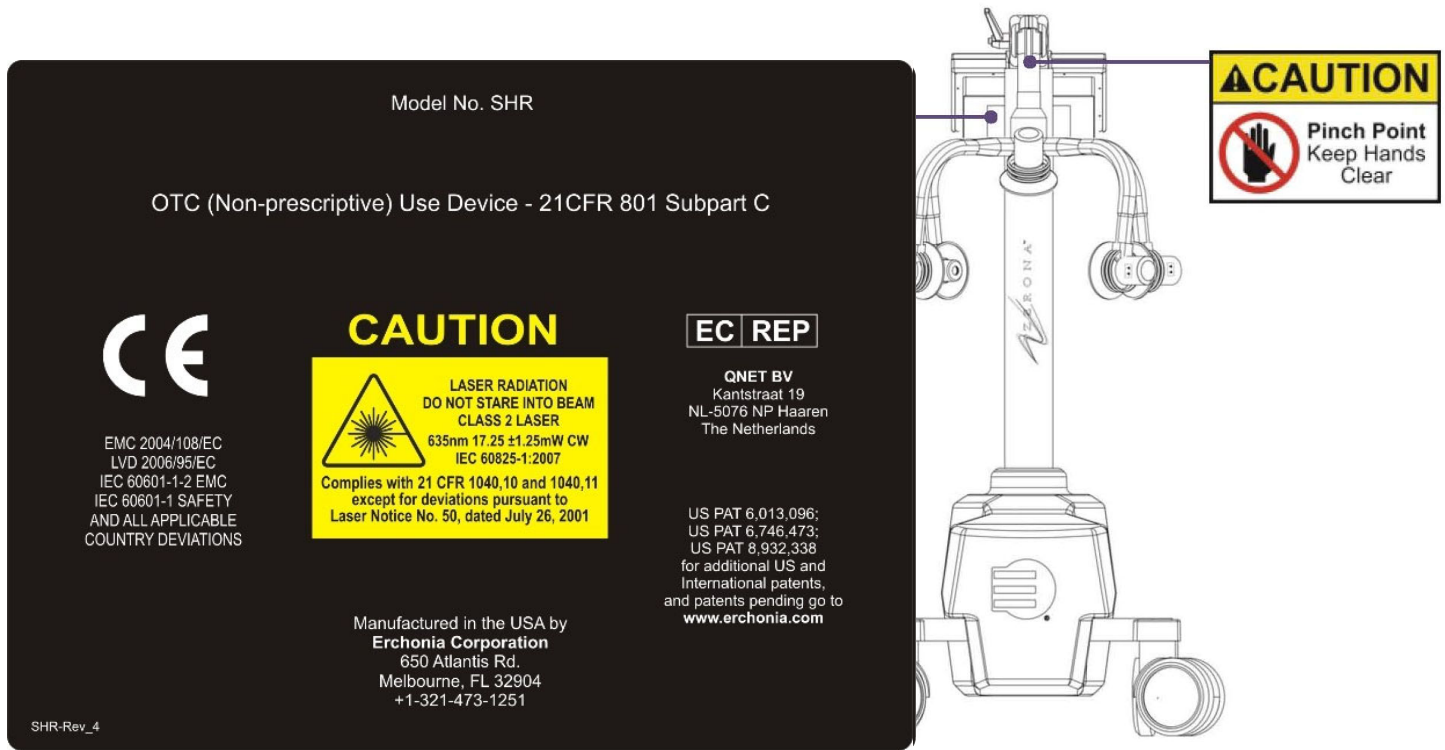
Inspect electrical cord and associated connectors before each use. **ONLY** use the power cord supplied with the device. Using a different power cord could damage the device

# LABELS

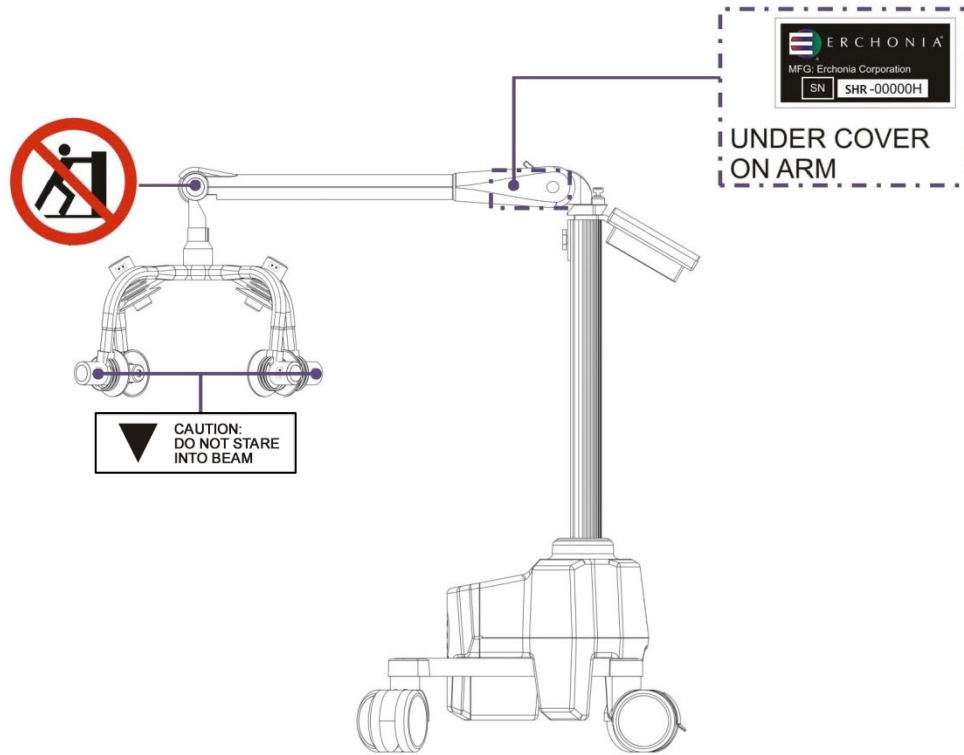
The device is manufactured in accordance to the Good Manufacturing Practices set forth by the FDA, ISO Standards (International) and CE (Conformité Européenne or European Conformity) standards and testing results per Article 9. The device is a Class I Shock Protection and a Class II Medical device. Each of these governing agencies requires specific labeling. All required labels are affixed per the relevant codes. Each label is pictured and described in this section. Additionally, the placement of each label on the device is communicated.

The following diagram shows the compliance labels and their placement.

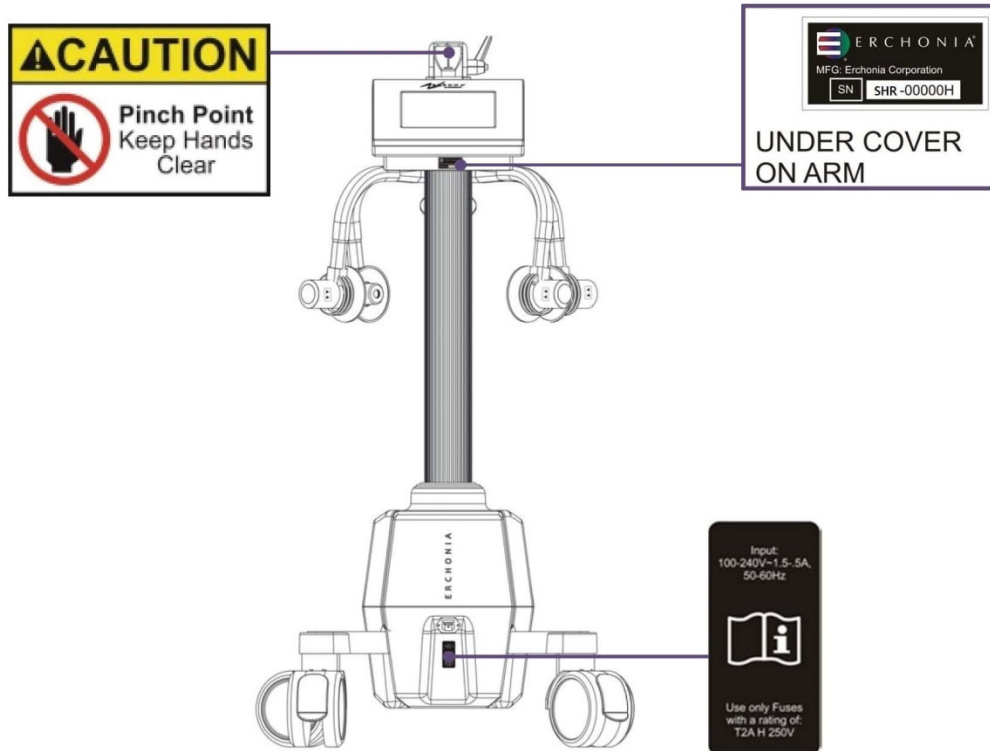
The large black background label is this primary label and is compliant to FDA and ISO standards; the image captures the FDA code regulated classifications and International criteria. One of the (2) identified pinch points is labeled below.



The following diagram shows the location of the device arm serial number, no pushing reminder and each of the laser diodes (6) have a label affixed that show the direction of the laser beam output.



The following diagram shows one of the (2) identified pinch points, the device base serial number and electrical input label containing Volts, Amps and fuse specifications.



## RISKS AND BENEFITS

Through clinical studies, the Zerona® Z6 OTC Laser has been shown to reduce the circumference of the waist, hips and thighs.

The risks and benefits of the Zerona® Z6 OTC laser are based on participant's feedback and all other clinical studies conducted using Erchonia low level lasers for circumference reduction.

### RISKS

There have been no side effects as a result of use of the Erchonia ZERONA® Z6 OTC laser. There have also been other research studies using Erchonia low level lasers. In these studies, no side effects resulted from or reported by any participant from use of the device.

The only known or anticipated risk with the use of the laser device is that long-term exposure to laser light could cause damage to eyesight. As a precaution, when you are giving the treatments with the Zerona® Z6 OTC, ensure that the client wears the special darkened protective glasses to block out the light.

Women who are pregnant or trying to get pregnant should not be treated.

The following conditions or medications may affect the client's outcome or weaken fat mobilization or metabolic pathways:

- Steroid medications (may cause weight or girth gain)
- Diabetes (may experience rapid weight or girth gain)
- Metabolic Disorders (may experience rapid weight or girth gain)
- Lymphatic Impairment (fat mobilization may be hindered)
  - Previous abdominoplasty, C-section, liposuction (procedures that have created scar tissue around the treatment site)

It is possible that the client will not get the circumference reduction that they desire.

# BENEFITS

In the past, Erchonia's low level laser technology was only available in a doctor's office for circumference reduction of the waist, hips and thighs. Erchonia first received FDA market clearance in 2010 as a prescription only device.

The application of Erchonia's low level laser technology has been proven through clinical trials to induce transitory pore formation, releasing collected fatty material, which is absorbed, degraded by the lymphatic system and removed from the treated area, creating a slimming outcome based upon the volume reduction of fat cells.

In 2008, Erchonia conducted a yearlong clinical trial study to determine the effectiveness of the Erchonia Zerona for non-invasive circumference reduction of the waist, hips and thighs with 6 treatments over a 2-week time frame.

A double-blind, randomized, multi-site, and placebo-controlled study with 67 participants was conducted in which patients on average, lost 3.50 inches/ 8.9 cm\* from their waist, hips and thighs. These results are dramatic compared to the placebo group, which only lost an average of half an inch.

In 2014, 22 participants between the ages of 18 to 65 participated in a multi-site clinical study that was conducted in the United States using the Zerona Z6 OTC device. Participants received six treatments over a two-week period. Throughout the study, participants were measured at the end of each week. The participants saw a combined circumference reduction of the waist, hips and thighs averaging 3.7 inches at the end of 2 weeks.

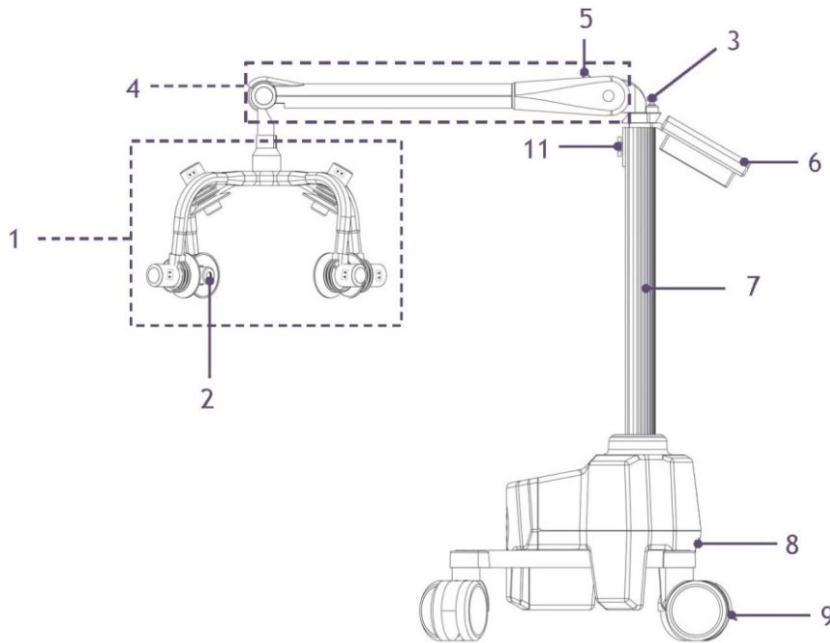
**To achieve the best results, you must carefully read and follow the instructions in this booklet.**

\*individual results may vary

# SECTION 2: ZERONA Z6 OTC OVERVIEW & ASSEMBLY

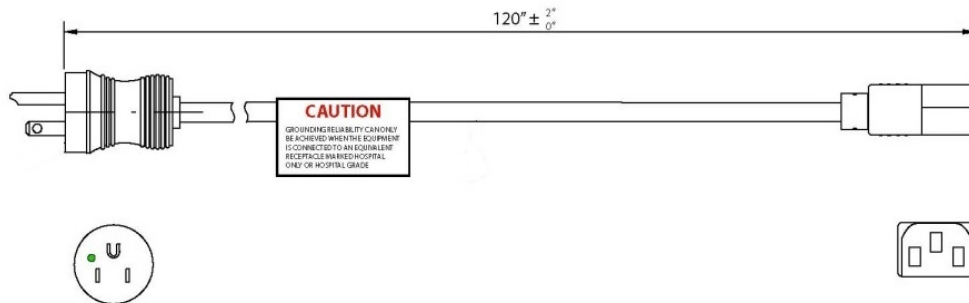
## ZERONA Z6 OTC ASSEMBLY

### DIAGRAM OF MACHINE



- 1. Laser Head Assembly
- 2. Laser Output Head
- 3. Power Safety Lockout Key
- 4. Laser Arm
- 5. Arm Lock
- 6. Touch Screen
- 7. Main Upright of Base
- 8. Power Inlet/Fuse Holder
- 9. Wheel Locks
- 10. Electrical Connector - See Fig. 2
- 11. Locking Knob

Fig. 1



12. Electrical Power Cord

# ASSEMBLY DIRECTIONS

- 4. Laser Arm
- 7. Main Upright of base
- 10. Connector & Cable
- 11. Locking Knob

- This may require two people.
- Remove the locking knob [11] from the main upright of base.
- Once the locking knob is removed, hold the laser head assembly and laser arm piece while you attach the base [7] and the arm piece [4].

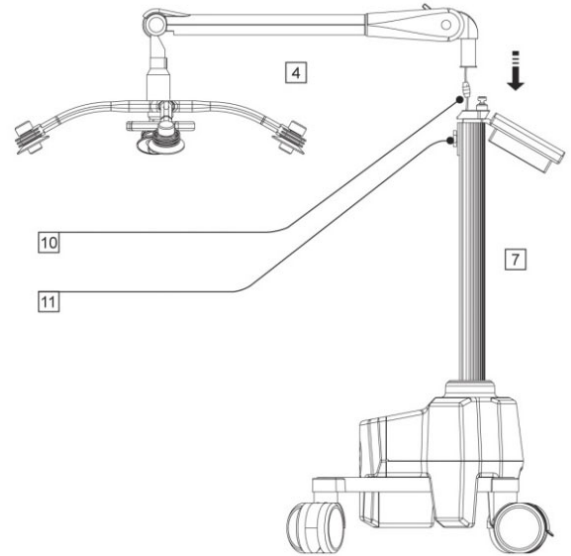
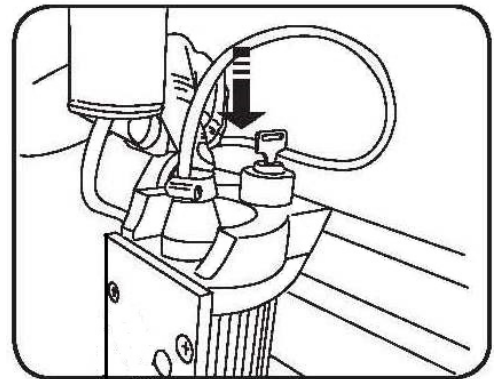
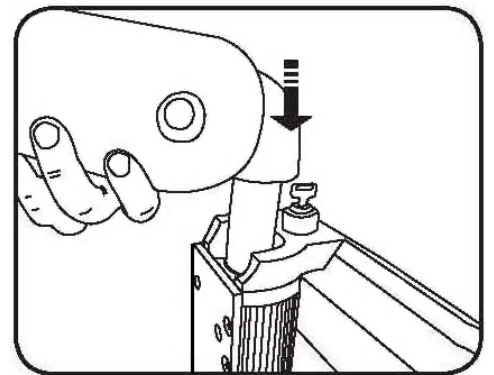


Fig. 2

- Gently feed the laser arm cable into the base device's upright shaft.

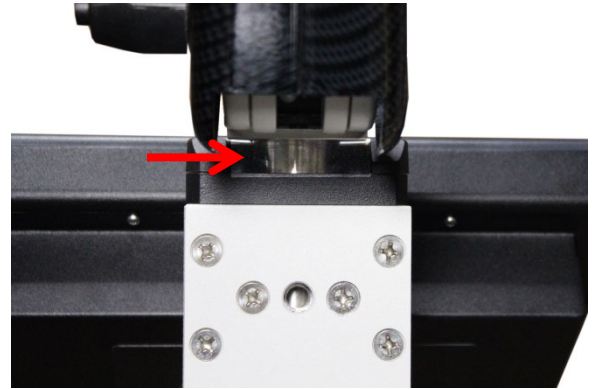


- With the cables fed into the base device's upright shaft, insert the arm tube into the base. Make sure the tube is aligned with the locking knob hole.



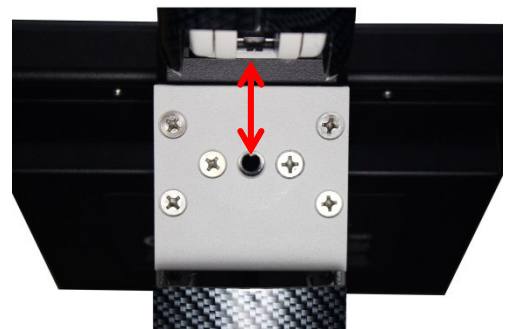
## ASSEMBLY DIRECTIONS (CONTINUED)

INCORRECT

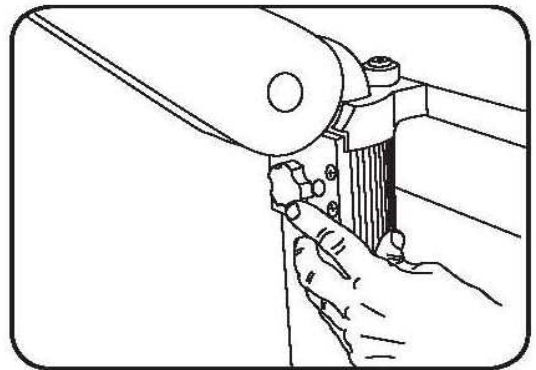


- Once the arm is inserted, gently push it down into the base until the two pieces are flush and the hole opening for the locking knob is aligned.

CORRECT



- With the holes aligned, thread the locking knob into place. This will keep the main head assembly from unwanted rotation during use.



- Access the laser arm cable by pulling the tab on the upright base cover as shown.
- Reach in and pull out the laser arm cable for ease of connecting.



## ASSEMBLY DIRECTIONS (CONTINUED)



### WARNING-SHOCK HAZARD

To avoid risk of electric shock, ensure that the electrical power cord is disconnected from electric supply when connecting electrical connector.

- Insert the two halves of the electrical connection [10] together by aligning the yellow markings slightly pushing the outlet into the inlet as shown.



- Twist the ring on the connector clockwise until it can no longer be tightened to make sure it is secure. This is important so that the two halves do not separate while within the device.
- Place the connector and cable inside the upright base shaft and secure the cover.



- Now that the laser has been put together, insert the electrical power cord [12] into the power inlet [8] located at touch screen side of the base device. See figure 1.
- Plug the electrical power cord [12] into any standard electrical outlet for your country/region. The device includes a transformer which converts AC power to match the power output (i.e. 110V or 240V). Only a 3-plug prong adaptor is required. Once the adaptor is fixed to the plug end, insert it into the wall socket. Disconnecting this cord while the machine is in operation will stop the machine, so this can be used as a second or emergency turnoff.



### WARNING-SHOCK HAZARD

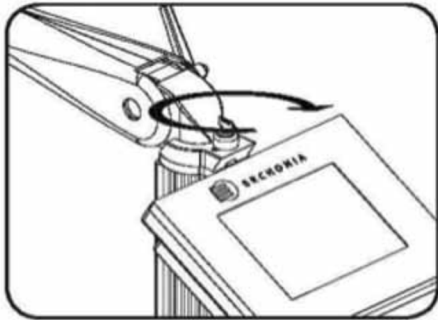
To avoid risk of electric shock, make certain that the device is grounded by connecting only to a grounded electrical socket (3-prong) conforming to the applicable national and local electrical codes.

## ASSEMBLY DIRECTIONS (CONTINUED)



**PRECAUTION - DO NOT** connect to an electrical outlet controlled by a wall switch or dimmer. For continued safety and performance, use only the power cord supplied by Erchonia. Do not position equipment so that in an emergency it is difficult to disconnect the power cord from the electric supply.

- Now insert one of the two provided keys into the keyhole on the base device.



The main power switch is the Power Safety Lockout Key switch on top of the touch screen. There is an external key for the switch to turn the device ON or OFF. (“O” = OFF and “|” = ON). In the OFF position, the device is locked, and the external key can be removed. This is a code-regulated feature to ensure no unauthorized use of the laser device. The device will not operate in the OFF position. Turning the key to the OFF position while using the device will immediately power off the device.

**NOTE:** It is highly recommended that you separate the two keys provided and store one of them in a safe place until needed.

## BACKUP BATTERIES

**ATTENTION:** The laser device has two internal back-up batteries that retain certain aspects of the device’s software memory in the case of power failure or shipping the device. These batteries receive a charge whenever the device is powered ON. It is important that the device is powered ON when first received by the user for a minimum of 2 minutes, as well as bi-weekly for a minimum of 2 minutes, to ensure proper charge of the batteries. Failure to do so may result in certain aspects of the memory being lost, which will require you to ship the device to the manufacture.

# SECTION 3: ZERONA Z6 OTC OPERATION

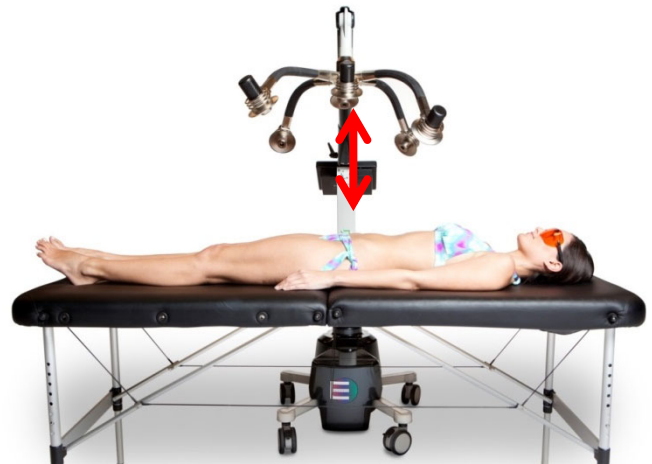
## BASICS AND ESSENTIALS

Now that you understand the basic features of the Zerona Z6 OTC laser, it is important to understand how to use them when administering treatments.

### MOVING THE LASER ARM

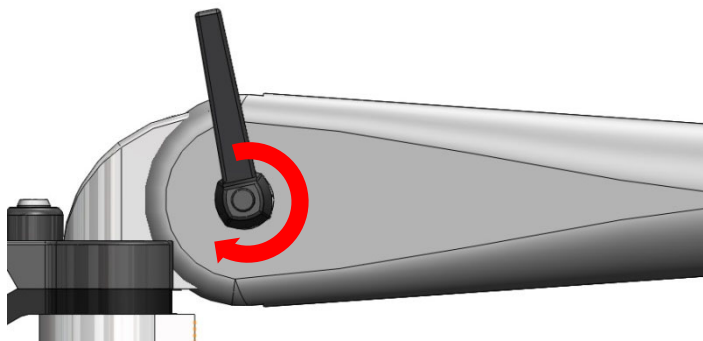
The arm is moved up and down using the laser arm, not the laser output head or flexible arms.

The arm tension has been set by the manufacture to a neutral state. This means that the arm will not move up or down unless you physically move it.



Once you have maneuvered the laser arm to the desired position, as a secondary precaution use the arm lock and apply a small amount of tension.

To do this, simply rotate the arm lock clockwise to increase the tension and counter-clockwise to decrease the tension.



### WHEEL LOCKS

The Zerona Z6 OTC laser device has four antistatic wheels to make moving it around easy. Once you have moved the laser to the desired location, be sure to engage the wheel locks to avoid excessive movement.





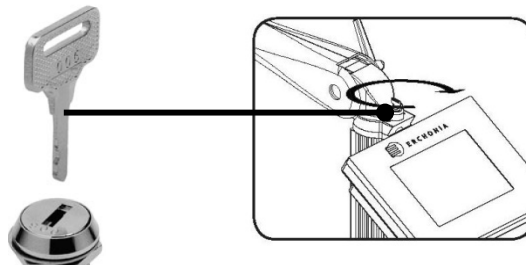
### WARNING-TIPPING HAZARD

When moving the device, ensure that the laser arm is positioned all the way down and secured in this position with the arm lock. Be careful to make sure the device does not tip while moving it.

## ZERONA Z6 TOUCH SCREEN

The touch screen is both a display screen that provides information to the user and an input panel that allows the user to operate the device by touching the appropriate icon.

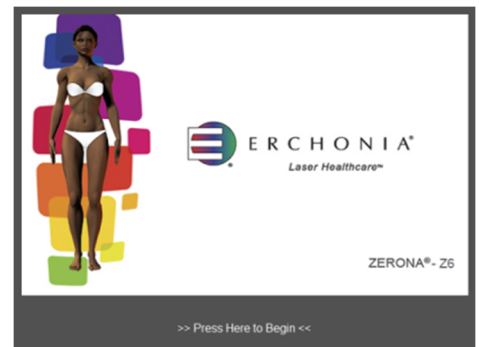
- Make sure the power is connected.
- Turn the device ON by placing one of the keys into the key lock and turning it clockwise to the ON “|” position.



- When the device is ready for use, after about 30-45 seconds, the touch screen will display the start screen.

- Touch anywhere on the “**Press Here to Begin**” bar which is found across the bottom of the screen. This will take you to the ZERONA Z6 Mode screen.

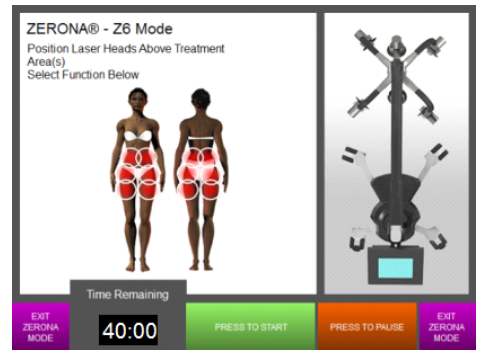
### Start Screen



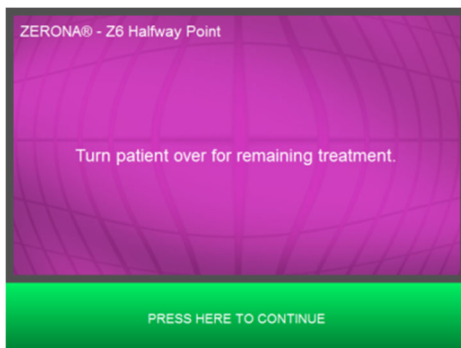
**PRECAUTION - DO NOT** use sharp objects such as a pencil point or ballpoint pen to operate the buttons on the touch screen as damage may result. Avoid using abrasives (including paper towels) on the touch screen display window.


- Touch **“PRESS TO START”** icon to begin the non-invasive procedure. Once you start the procedure you will observe the following:
  - six visible red laser diodes turn on rotating clockwise;
  - the **“Time Remaining”** window starts counting down;
  - the **“PRESS TO START”** icon will flash blue and grey, and the text will display **“PRESS TO STOP”**.
- To pause, touch the **“PRESS TO PAUSE”** icon. When the procedure is paused, you will observe the following:
  - six red laser diodes and rotation turn off;
  - the time remaining counter pauses;
  - the **“PRESS TO PAUSE”** icon will flash dark green and light green, and the text will display **“PRESS TO RESUME”**;
- To resume the procedure, touch the flashing **“PRESS TO RESUME”** icon.

### Zerona Z6 Mode Screen



### Zerona Z6 Halfway Point Screen



Once the **“Time Remaining”** display reaches 20:00 the lasers will power off,  you will hear 3 audio “beeps”, and the **“ZERONA Z6 Halfway Point”** screen will display.

**NOTE:** The device is preprogrammed to treat for a total of 40 minutes.

After the first 20 minutes of treatment, the device will pause, displaying the **“Zerona Z6 Halfway Point”** screen. This is to give the client time to change from the front to the back treatment position.

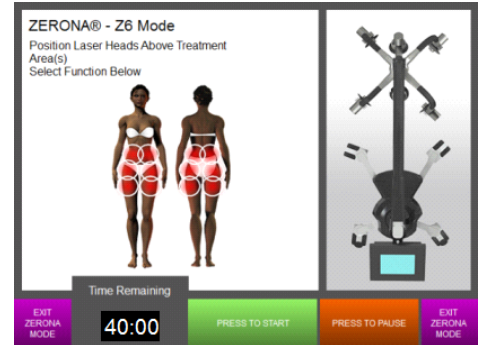
Once the client is ready in the back treatment position (lying on their stomach) and you have properly positioned the diodes, touch **“PRESS HERE TO CONTINUE”** icon to complete the second 20 minutes of the Zerona Z6 treatment procedure.

# ZERONA Z6 TOUCH SCREEN (CONTINUED)

Located on the bottom are the active command icons. Starting from left to right the icons are:

- **“EXIT ZERONA MODE”** this will return you to the start screen.
- Located to the right of the **“Exit Zerona Mode”** icon is the **“Time Remaining”** display that shows the current treatment time remaining. The countdown display activates once the treatment is started.
- Next to the Time Remaining display is the **“PRESS TO START”** icon; this will start and stop the treatment program.
- Next is the **“PRESS TO PAUSE”** icon, when touched, this icon will change colors and flash **“PRESS TO RESUME”** allowing the client to take a break without resetting the time remaining. Touch the **“PRESS TO RESUME”** icon to continue with treatment.

Zerona Z6 Mode Screen



## DEVICE INFORMATION SCREEN

To access the **“Device Information”** screen touch the Erchonia **“E logo”** on the start screen. This multi-functional display screen shows the following:

- Manufacturer’s contact information;
- General information for your device including software revision, device computer date which is the date based on central standard time USA, device serial number and total number of treatments the device has operated;
- On the upper right is a Green circle with the word OK. This is the device RE-SET button and is only to be used at the instruction of an Erchonia Manufacture representative.

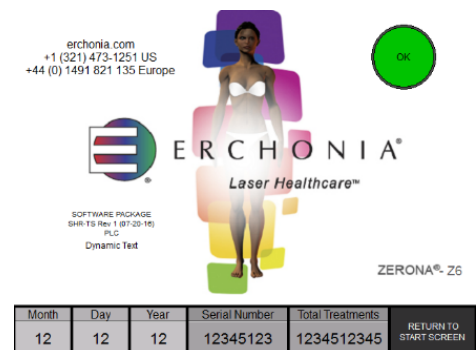
### Manufacturer’s Information

Erchonia Corporation  
 Telephone: 1-321-473-1251  
 Email: info@erchonia.com

Start Screen



Device Information Screen



# BASICS AND ESSENTIALS (CONTINUED)

## PROTECTIVE EYEWEAR

- The Zerona Z6 OTC is classified by the FDA/IEC as a Class 2 laser device. This represents a current standard for use to ensure the safety of the client.
- A Class 2 laser is determined to have a chronic viewing hazard. Pointing the laser beam directly into the eye and maintaining it there for an extended period could prove to be damaging.
- To ensure there is no possible instance of residual effect, we have included a pair of specialty glasses for use by the client during treatment. The safety glasses, sufficiently and effectively block the laser light spectrum at OD 2+ @ 635nm, OD 0.75 @ 405nm VLT60.

Height: 40 mm

Width: 145 mm

Length: 165 mm



**WARNING-** The client should always be correctly fitted with the safety glasses provided before turning on the laser and doing any treatment.

# PROPER DIODE PLACEMENT

Have your client put on the safety glasses and lay on their back.

Position the center laser diodes and adjustable laser arms, no greater than 3-4 inches / 9 cm away from the client's target treatment areas. The goal is to maximize the coverage of the laser beams to the target treatment areas of fat.

While the client is lying on their back you should check for the areas of the greatest accumulation of fat, which will tend to be on the sides of the waist, hips, and thighs.

Using the center laser diodes as your guide, place the center diodes slightly below the navel. Position the adjustable laser arms so they are targeting the greatest accumulation of fat on the left and right side of the waist, hips, and thighs. **NOTE:** It is important to align the diodes so that the line-generated beam reaches the skin's surface at 90° or perpendicular.

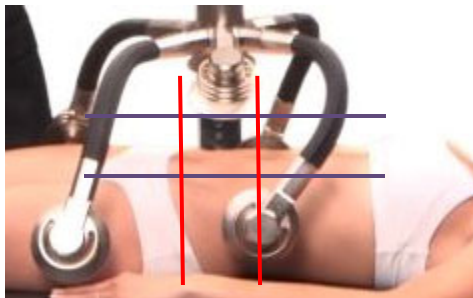


The following image shows proper laser diode placement to effectively treat the fat pockets of the waist, hips and thighs.

- The center diodes are positioned over the area of greatest midline fat accumulation.
- The lateral (side) adjustable laser heads are positioned over the areas of greatest waist and thigh fat accumulation on either side of the body.

## Anterior (Front) Treatment Diode Placement

Perpendicular Placement of Diode in Reference to the Treatment Area



# PROPER DIODE PLACEMENT (CONTINUED)

## LINE-GENERATED BEAM PARAMETERS 3.0 Inch (7.5 cm) Line-Generated Beams




## 6 inches (15 cm) separating adjacent Line-Generated Beams



Use the palm of your hand to position the adjustable laser heads, shifting the flexible arms to the region of greatest fatty accumulation, as shown.



Now that you've positioned the diodes properly in the targeted areas, press the start button to begin. With the lasers on, check the positioning of the heads for coverage of targeted areas. We suggest dimming or turning off the lights to ensure that you have maximized the coverage area. When the first 20 minutes are up, the lasers will pause and  3 audio beeps will emit from the machine.

Raise the laser arm from the client, and have them roll over and lie on their stomach. Lower the laser arm over the client, positioning the center laser diodes in line with an identical position as if the client were lying on their back (approx. 1-2 inches / 5 cm below the belly button).

Position the center laser diodes and adjustable laser arms, no greater than 3-4 inches / 9 cm away from the client's target treatment areas. The goal is to maximize the coverage of the laser beams to the target treatment areas of fat.

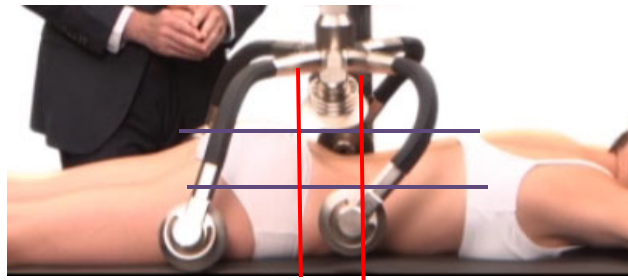
Position the adjustable laser arms so they are targeting the greatest accumulation of fat on the left and right side of the waist, hips, and thighs. **NOTE:** It is important to align the diodes so that the line-generated beam reaches the skin's surface at 90° or perpendicular.

The following image shows proper laser diode placement to effectively treat the fat pockets of the waist, hips and thighs.


- The center diodes are positioned over the area of greatest midline fat accumulation.
- The lateral (side) adjustable laser heads are positioned over the areas of greatest waist and thigh fat accumulation on either side of the body.

## Posterior (Back) Treatment Diode Placement

### Perpendicular Placement of Diode in Reference to the Treatment Area



Touch the start button again and check the positioning of the line generated beams to ensure proper positioning.



After the second 20-minute posterior (back) treatment, the laser will automatically turn off and alert you with  3 audio beeps.

Raise the laser arms from the client. The client can get dressed, leave the office, and resume their daily lives.

# ZERONA®-Z6 OTC TREATMENT PROTOCOL

Now that you understand how to operate the Zerona Z6 OTC device and how to position the diodes properly, the following section describes the steps to follow each time you administer a treatment with the device to a client. You will need to administer six treatments with the Zerona Z6 OTC device to the client over two weeks. This means three treatments each week, with each treatment every other day.

Follow the treatment plan example for best results

- You must always begin by having the client put on the safety glasses.
- Have the client lie comfortably flat on their back on the treatment table so that the front of their body is facing upwards.
- Position the laser diodes properly over the client's targeted area following the procedure described in the previous section of this booklet.
- Once the diodes are positioned properly in the targeted areas, touch the **START** icon. With the lasers on, check the positioning of the heads for coverage of targeted areas as described in the previous section of this booklet.
- When the first 20 minutes of treatment on the client's front body area is done, the lasers will pause and the device will alert you with  3 audio beeps.
- Raise the laser arm above the client and have the client roll over to lie comfortably on their stomach.
- Lower the laser arm and position the laser diodes properly over the backside of the client's targeted area following the procedure described in the previous section of this booklet.
- Once the diodes are positioned properly in the targeted areas, touch the **START** icon again. With the lasers on, check the positioning of the heads for coverage of targeted areas as described in the previous section of this booklet.
- After the second 20 minutes of treatment for the client's back body area is done, the laser will automatically turn off and alert you with  3 audio beeps.
- Raise the laser arm off the client.
- The client may get up from the treatment table and remove the safety glasses. Their treatment procedure is done.
- Turn the key to the OFF position.

# ZERONA®-Z6 OTC CLIENT QUALIFICATION CHECKLIST

You should always use the Zerona Z6 OTC Client Qualification Checklist to make sure the client is suitable for treatments before starting any treatments.

**A. CLIENT QUESTIONS:** Ask the client all four questions below and check the answer as 'Yes' or 'No'

Question	Yes	No
Are you under 18 years of age?	<input type="checkbox"/>	<input type="checkbox"/>
(Female clients only): Are you pregnant or do you think you might be pregnant? <input type="checkbox"/> Not applicable, client is male	<input type="checkbox"/>	<input type="checkbox"/>
Do you have any open wounds (sores, cuts, ulcers, etc.)?	<input type="checkbox"/>	<input type="checkbox"/>
Do you have or are you being treated for any cancerous growths?	<input type="checkbox"/>	<input type="checkbox"/>

Check the correct box below and follow the action

<input type="checkbox"/>	All the 'No' boxes are checked => <b>continue with the client qualification checklist</b>
<input type="checkbox"/>	One or more 'Yes' boxes is checked => <b>the client does not qualify for Zerona Z6 OTC</b>

## B. BODY MASS INDEX (BMI)

Calculate the client's Body Mass Index (BMI) by following these steps:

1. Measure the client's height in inches and write it in the table below
2. Measure the client's weight in pounds and write it in the table below
3. Work out the client's Body Mass Index (BMI) using the Body Mass Index Table on the next page by following these steps:
  - a. Find the client's height in inches that you measured in the furthest left-hand side column of the table.
  - b. Move your finger from left to right across the row for that height until you find the client's weight in pounds that you measured. This number may fall in between two other numbers. Move your finger to the closest number, which may be the number above or below the client's weight number.
  - c. Move your finger up the column for that weight until you reach the top row labeled 'BMI.' The number you land on is the client's BMI.

For example, a client whose height is 62 inches and weight is 160 pounds (closest to 158 pounds in that height row) has a BMI of 29.

4. Write the client's BMI in the table below.

Client's height (inches)	
Client's weight (pounds)	
<b>Client's Body Mass Index (BMI)</b>	

Check the correct box below and follow the action

<input type="checkbox"/>	BMI is 30 or less => <b>the client qualifies for Zerona Z6 OTC treatment.</b>
<input type="checkbox"/>	BMI is more than 30 => <b>the client <u>does not</u> qualify for Zerona Z6 OTC</b>

## Body Mass Index Table

	Normal										Overweight										Obese										Extreme Obesity									
BMI	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54				
Height (inches)	Body Weight (pounds)																																							
58	91	96	100	105	110	115	119	124	129	134	138	143	148	153	158	162	167	172	177	181	186	191	196	201	205	210	215	220	224	229	234	239	244	248	253	258				
59	94	99	104	109	114	119	124	128	133	138	143	148	153	158	163	168	173	178	183	188	193	198	203	208	212	217	222	227	232	237	242	247	252	257	262	267				
60	97	102	107	112	118	123	128	133	138	143	148	153	158	163	168	174	179	184	189	194	199	204	209	215	220	225	230	235	240	245	250	255	261	266	271	276				
61	100	106	111	116	122	127	132	137	143	148	153	158	164	169	174	180	185	190	195	201	206	211	217	222	227	232	238	243	248	254	259	264	269	275	280	285				
62	104	109	115	120	126	131	136	142	147	153	158	164	169	175	180	186	191	196	202	207	213	218	224	229	235	240	246	251	256	262	267	273	278	284	289	295				
63	107	113	118	124	130	135	141	146	152	158	163	169	175	180	186	191	197	203	208	214	220	225	231	237	242	248	254	259	265	270	278	282	287	293	299	304				
64	110	116	122	128	134	140	145	151	157	163	169	174	180	186	192	197	204	209	215	221	227	232	238	244	250	256	262	267	273	279	285	291	296	302	308	314				
65	114	120	126	132	138	144	150	156	162	168	174	180	186	192	198	204	210	216	222	228	234	240	246	252	258	264	270	276	282	288	294	300	306	312	318	324				
66	118	124	130	136	142	148	155	161	167	173	179	186	192	198	204	210	216	223	229	235	241	247	253	260	266	272	278	284	291	297	303	309	315	322	328	334				
67	121	127	134	140	146	153	159	166	172	178	185	191	198	204	211	217	223	230	236	242	249	255	261	268	274	280	287	293	299	306	312	319	325	331	338	344				
68	125	131	138	144	151	158	164	171	177	184	190	197	203	210	216	223	230	236	243	249	256	262	269	276	282	289	295	302	308	315	322	328	335	341	348	354				
69	128	135	142	149	155	162	169	176	182	189	196	203	209	216	223	230	236	243	250	257	263	270	277	284	291	297	304	311	318	324	331	338	345	351	358	365				
70	132	139	146	153	160	167	174	181	188	195	202	209	216	222	229	236	243	250	257	264	271	278	285	292	299	306	313	320	327	334	341	348	355	362	369	376				
71	136	143	150	157	165	172	179	186	193	200	208	215	222	229	236	243	250	257	265	272	279	286	293	301	308	315	322	329	338	343	351	358	365	372	379	386				
72	140	147	154	162	169	177	184	191	199	206	213	221	228	235	242	250	258	265	272	279	287	294	302	309	316	324	331	338	346	353	361	368	375	383	390	397				
73	144	151	159	166	174	182	189	197	204	212	219	227	235	242	250	257	265	272	280	288	295	302	310	318	325	333	340	348	355	363	371	378	386	393	401	408				
74	148	155	163	171	179	186	194	202	210	218	225	233	241	249	256	264	272	280	287	295	303	311	319	326	334	342	350	358	365	373	381	389	396	404	412	420				
75	152	160	168	176	184	192	200	208	216	224	232	240	248	256	264	272	279	287	295	303	311	319	327	335	343	351	359	367	375	383	391	399	407	415	423	431				
76	156	164	172	180	189	197	205	213	221	230	238	246	254	263	271	279	287	295	304	312	320	328	336	344	353	361	369	377	385	394	402	410	418	426	435	443				

Source: Adapted from Clinical Guidelines on the Identification, Evaluation, and Treatment of Overweight and Obesity in Adults: The Evidence Report.

# MEASUREMENT PROTOCOL

➤ We recommend taking client measurements at the following times:

1. Before the first treatment
2. After the 6th treatment

- To take proper measurements, the client should stand with their feet shoulder width apart. Have them relax their muscles. Do not let them “suck it in,” as this is a typical reflex of most clients.
- When taking measurements, make sure that the tape is pulled just tight enough so that it isn't sagging but at the same time it is not too snug.
- You may also want to take before and after photos of your clients. These are both great ways to show your client their progress!



# SECTION 4: MAINTENANCE & WARRANTY INFORMATION

## TECHNICAL INFORMATION

Technical documentation required by the customer, in case of necessary reparations, will be provided by Erchonia in the US and our EU agent, internationally. These documents will be supplied once the manufacturer makes the determination that the requested documents do not constitute a disclosure of proprietary or patent protected information and are a part of the filed and documented technical file.

## SERVICE AND REPAIR

If a device requires service, contact the Erchonia Service and Repair Department at:

Telephone: +1-321-473-1251

When requesting service or repair, please provide the following information to the service representative:

- Device serial number (located on the back label)
- Description of the problem
- Name of the person to contact

## RETURNING A DEVICE FOR SERVICE

- Before sending a device to the Erchonia Service and Repair Department for repair, obtain a service order (SO) number from the service representative.
- Pack the device in the original containers (if available) or equivalent packaging. Be sure the assigned service order number appears on the package.

### **Return the Device to:**

Erchonia Corporation

650 Atlantis Rd.

Melbourne, FL. 32904 USA

Attention: Service and Repair Department (*SO Number*)

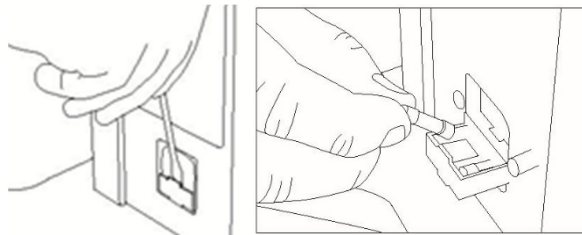
**NOTE:** For international customers, PRIOR to sending a device in for repair you must obtain from the Erchonia Service department an annually revised FDA Form 2877. The Radiation Control form (2877) will be sent to you partially complete, containing regulatory information. To complete, fill in the unique information associated with your device and the shipment thereof; such as serial number, port, etc. The completed form 2877 must accompany your shipment affixed to the outside of the package. Failure to include the form in the shipment may result in customs delays and fines. Any resulting fines are the responsibility of the customer.

# MAINTENANCE AND CLEANING

## *MAINTENANCE*

The Zerona Z6 OTC, if used per the instructions contained within this manual, will operate efficiently for years. To ensure proper care, it is advisable for you to perform the following:

1. Regular visual inspections to ensure there is no external damage other than normal wear and tear. Inspect all cords for signs of excessive wear (cuts in insulation or fraying). If during these inspections, you identify an area of concern, please contact Erchonia Corporation to determine if action is required.
2. If you notice a change in the performance of the device, please contact Erchonia Corporation to determine if action is required.
3. The internal components should not require any maintenance, however if an issue arises, which will show itself in the form of altered performance, the device must be sent to Erchonia Corporation.
4. Since the device contains a touchscreen interface, periodic cleaning of the touchscreen will be necessary. To clean the touchscreen, use a nearly dry cloth containing one of the mild cleaning agents listed below. Ensure there is **NOT** an excess of fluid.
5. The touchscreen back up battery must be replaced every five years. This must be done by manufacturer.
6. If the fuses require replacement, unplug the power cord from plug source, locate power inlet / fuse holder (figure 1 item 9), pull fuse holder out of enclosure, replace the two fuses (**T2AH 250V Fuses only**) and reinsert.



## *CLEANING INSTRUCTIONS*

### *ROUTINE USER MAINTENANCE*

Clean the external surfaces of the device	Weekly, or as needed
Clean the touch screen display	Weekly, or as needed
Clean laser optics	Weekly, or as needed
Clean Laser glasses	Between each patient use
Connect power to the device and turn the key to the ON position, powering the device ON. Keep device powered ON for a minimum of 2 minutes. This is to properly maintain the charge in backup batteries. Failure to do so may result in certain aspects of the memory being lost.	When the device is first received by the user and bi-weekly

## ***CLEAN THE EXTERNAL SURFACES OF THE DEVICE***

Use a cloth dampened with non-caustic cleaning solution, such as mild soap and water, isopropyl alcohol, or a “hospital-grade” disinfectant, to wipe the external surfaces of the device. Dry with a clean cloth, or allow to air dry.



**DO NOT** permit any foreign materials or liquids to enter the device. Take care to prevent any foreign materials including, but not limited to, inflammables, water, and metallic objects from entering the device. These may cause device damage, malfunction, electrical shock, fire, or personal injury. To achieve the specified level of protection against spilled or splashed liquids, unplug the device from the power supply and thoroughly dry all exposed surfaces of this device and allow to dry thoroughly prior to operation.

## ***CLEAN THE TOUCH SCREEN DISPLAY***

Apply an alcohol-based cleaner to a soft cloth to clean the touch screen display.



**DO NOT** spray or pour cleaning agents directly on the device or touch screen. You may damage the console, touchscreen, and the system electronics.

## ***LASER OPTICS CLEANING***

Use lens paper or lens cloth **ONLY**. Abrasive material could cause laser light beam fragmentation, which may reduce the effectiveness of the treatment.

## ***LASER GLASSES CLEANING***

Use only mild soap

# ***STORAGE***

The Zerona Z6 OTC laser should be stored in a cool, dry place out of the reach of children.

# WARRANTY

## *LIMITED WARRANTY*

The Erchonia® laser device is warranted to be free from defects in material and workmanship for a period of TWO YEARS from the date of purchase.

For your device to be processed through the Service and Repair Department efficiently, contact the department prior to submitting your product. Repairs and Warranty work NOT coordinated through the Service and Repair Department prior to receipt can be delayed.

Items being sent in from overseas require special paperwork, which is available through the Service and Repair department. If paperwork is not included your package will be delayed by customs.

## *TERMS AND CONDITIONS*

- Shipping required to facilitate warranty repair and or maintenance issues within the first 90-days, will be paid by manufacturer.
- Shipping required to facilitate warranty repair and or maintenance issues after 90-days, is the financial responsibility of the customer.
- Warranties of Erchonia Corporation products are not transferable unless sold by a company-approved distributor, reseller and/or leasing company.
- The warranty DOES NOT cover instances involving or damages resulting from:
  - Accident, misuse or abuse
  - Lack of responsible care
  - Alteration or disassembly
  - Loss of parts
  - Exposure to the elements
  - Ingress of liquid
  - Exposure to excessive electromagnetic frequency

## *PRODUCT CHANGES*

Erchonia Corporation reserves the right to make changes and improvements to its products without incurring any obligation to incorporate such improvements in products previously sold.

## CONTACT US

Questions? If for any reason you are dissatisfied with this product, warranty concerns or questions regarding proper operation, please contact our Erchonia Customer Care representative for assistance. Contact us at:

**Erchonia Customer Care**

**Phone: +1-321-473-1251**

**Email: [info@erchonia.com](mailto:info@erchonia.com)**

Or visit [erchonia.com](http://erchonia.com)

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Protected by US Patents #s 6,013,096; 6,746,473 and 8,932,383. For additional US and International patents and patent pending go to [www.erchonia.com](http://www.erchonia.com)

# TROUBLESHOOTING

Use the following information to help solve problems that may arise during use. If you still have questions, please call Erchonia Customer Care at 1-321-473-1251, and we will be happy to help.

Problem	Possible Cause	How to Resolve
<b>POWER</b> Device does not power on	AC power cord not plugged into device or line power.	Connect power cord.
	Blown fuse.	Replace using the correct fuses.
	Lockout key not positioned correctly.	("O" = OFF and " " = ON)
	Blown fuse or tripped circuit breaker in building.	Contact a qualified service technician.
Diode(s) does not power on	Electrical connector came apart.	Remove head assembly from base assembly and reconnect connector. (See Assembly Directions Step 1)
Motors do not power on	Electrical connector came apart.	Remove head assembly from base assembly and reconnect connector. (See Assembly Directions Step 1)
Device protocol(s) time remaining, serial number, total treatments and credits remaining all display "0". Does not allow initiation of protocol(s)	Device backup batteries were not properly maintained	Contact Erchonia service department.

# GUIDANCE AND MANUFACTURER'S DECLARATION- ELECTROMAGNETIC EMISSIONS & IMMUNITY

Medical electrical equipment requires special precautions regarding EMC and must be installed and put into service according to EMC information provided in this document. Portable and mobile RF communications equipment can affect the medical electrical equipment.  
This declaration currently applies to the Zerona Z6 OTC device.

## GUIDANCE AND MANUFACTURER'S DECLARATION-ELECTROMAGNETIC EMISSIONS

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


Emissions test	Compliance	Electromagnetic environment-guidance
RF emissions CISPR 11	Group 1	The device uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class A	The device is suitable for use in all establishments, including domestic establishments and those directly connected to the public low voltage power supply network that supplies buildings used for domestic purposes.
Harmonic Emissions IEC 61000-3-2	Class B	
Voltage Fluctuations/Flicker Emissions IEC 61000-3-3	Complies	

### WARNING

- This device should not be used adjacent to other equipment. If adjacent use is necessary, the device should be observed to verify normal operation in the configuration in which it will be used.
- The use of accessories other than those specified for the device is not recommended. They may result in increased emissions or decreased immunity of the device.

# GUIDANCE AND MANUFACTURER'S DECLARATION-ELECTROMAGNETIC IMMUNITY

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

Immunity test	IEC60601-1-2 test level	Compliance level	Electromagnetic environment-guidance
<b>Electrostatic discharge (ESD)</b> IEC 61000-4-2	± 6 kV contact ± 8 kV air	± 6 kV contact ± 8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30 %.
<b>Electrical fast transient/burst</b> IEC 61000-4-4	± 2 kV for power supply lines ± 1 kV for input/output lines	± 2 kV for power supply lines ± 1 kV for input/output lines	Mains power quality should be that of a typical commercial or hospital environment.
<b>Surge</b> IEC 61000-4-5	± 1 kV line(s) to line(s) ± 2 kV line(s) to earth	± 1 kV differential mode ± 2 kV common mode	Mains power quality should be that of a typical commercial or hospital environment.
<b>Voltage dips, short interruptions and voltage variations on power supply input lines</b> IEC 61000-4-11	<5% UT (>95% dip in UT) for 0.5 cycle 40% UT (60% dip in UT) for 5 cycles 70% UT (30% dip in UT) for 25 cycles <5% UT (>95% dip in UT) for 5 sec	<5% UT (>95% dip in UT) for 0.5 cycle 40% UT (60% dip in UT) for 5 cycles 70% UT (30% dip in UT) for 25 cycles <5% UT (>95% dip in UT) for 5 sec	Mains power quality should be that of a typical commercial or hospital environment. If the user of the device requires continued operation during power mains interruptions, it is recommended that the device be powered from an uninterruptible power supply.
<b>Power frequency (50/60 Hz) magnetic field</b> IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.  Portable and mobile RF communications equipment should be used no closer to any part of the device including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.  <b>Recommended separation distance</b>
<b>Conducted RF</b> IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	3 Vrms	$d = 1.2/P$
<b>Radiated RF</b> IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz	3 V/m	$d = 1.2/P$ 80 MHz to 800 MHz
			$d = 2.3/P$ 800 MHz to 2.5 GHz  where P is the maximum output power rating of the transmitter in watts (W) per the transmitter manufacturer and d is the recommended separation distance in meters (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, <sup>a</sup> should be less than the compliance level in each frequency range. <sup>b</sup> Interference may occur near equipment marked with the following symbol: 

<sup>a</sup> Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy.

To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the Zeron Z6 OTC device is used exceeds the applicable RF compliance level above, the device should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the device.

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

**Notes:**

- UT is the AC mains voltage prior to application of the test level.
- At 80 MHz and 800 MHz, the higher frequency range applies.
- These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

**RECOMMENDED SEPARATION DISTANCES BETWEEN PORTABLE AND MOBILE RF COMMUNICATIONS EQUIPMENT AND THE DEVICE.**

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

Rated maximum output power of transmitter (W)	Separation distance according to frequency of transmitter (m)		
	150 kHz to 80 MHz $d = 1.2/\sqrt{P}$	80 MHz to 800 MHz $d = 1.2/\sqrt{P}$	800 MHz to 2.5 GHz $d = 2.3/\sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

For transmitters rated at a maximum output power not listed above, the recommended separation distance  $d$  in meters (m) can be estimated using the equation applicable to the frequency of the transmitter, where  $P$  is the maximum output power rating of the transmitter in watts (W) per the transmitter manufacturer.

**Notes:**

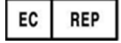
- At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.
- These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

# TECHNICAL SPECIFICATIONS

## Relevant Information



Erchonia Corporation  
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Melbourne, FL. 32904 USA



QNET BV  
Kantstraat 19  
NL-5076 NP Haaren  
The Netherlands



Trade name: Zerona® Z6-OTC  
Model Number: SHR

## Classifications

### US

- Device Class: II
- Laser Class: II

### UL/CSA

- Electrical Class 1

## Applicable Codes

### FDA

- 21CFR 820 - Quality System Regulations
- 21CFR 1040.10 and 1040.11 by laser Notice 50

### ISO

- 13485 - Medical Device Quality
- 60825-1 - Laser Safety

EMC 2004/108/EC

LVD 2006/95/EC

IEC 60601-1-2 EMC

IEC 60601-1-1 Safety

CB Certified

## Specifications

### Device

- Weight: 71lbs / 32kg
- Height: 70 in / 177.8 cm (Average - Adjustable)
- Full Color Touch Screen Control Center
- Four independent Adjustable Arms
- 4 locking Anti-static casters
- Powder coated metal frame
- Non-allergenic material finished with black carbon fiber
- Applied Part: Type B
- Leakage Current 0.3 -0.5  $\mu$ A (Micro Amps)

### Laser

- 6 line generating diode modules
- Output: 17.25mW +/- 1.25mW
- Wavelength: 635 nm +/- 10nm
- Duty Cycle: CW (Continuous 100%)

### Power

- Source: 100-240VAC 50-60Hz
- Fuse Rating: T2AH 250V

### Temperature

- Operating Temp: 59 to 85° F (15 to 29° C) <50% Relative Humidity
- Transporting: -22 to 158° F (-30 to 70° C) <75% Relative Humidity

### Atmospheric Pressure (Altitude)

- Rated operating altitude  $\leq$  2000 m
- Normal barometric pressure 80.0 kPa

## Legend:

FDA - US Food & Drug Administration, which includes the CDRH (Center for Device Radiological Health)

ISO - International Standards, Harmonized with US, Canadian, European and Asian standards

Doc No.	Issue Date	CR No.	Rev. Level	Rev. Date
ZERONA Z6-OTC-NC	3/23/2015	20-16-ADM	2	06/14/2016
ZERONA Z6-OTC-NC	3/23/2015	38-16-ADM	3	09/01/2016
ZERONA Z6-OTC-NC	3/23/2015	05-17-ENG	4	03/31/2017

Congratulations on completing the ZERONA Z6 OTC  
Installation and Proper Use Reference Guide.

You are now ready to provide great outcomes  
with the ZERONA Z6 OTC.



Non-Prescriptive/Non-invasive Fat Loss Laser

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