




SL3 User's Manual

This manual must be read thoroughly and understood prior to using the laser system. **CAUTION: Using the controls and adjustments or performing laser procedures, other than as specified herein, may result in hazardous radiation exposure.**

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Section 1: Introduction

INTRODUCTION

The SL3™ is a Class IV soft-tissue diode laser that can be used for a wide variety of soft-tissue procedures. This device uses a Gallium, Aluminum, Arsenic (GaAlAs) diode for the active medium, producing laser energy at 810 nanometer wavelength. Laser technology has been steadily evolving, allowing doctors to provide less invasive treatment for many dental procedures, both preventative and restorative. For a list of approved procedures please refer to page 12.

All of the safety features, warnings and precautions should be observed for the SL3™. All of the Safety Warning Labels are described in detail on page 15 of this guide. The SL3™ is especially designed to be compact, portable, reliable and user friendly. It provides the practitioner with a versatile instrument for applications ranging from excisions to vaporization of tissues. The diode laser energy is delivered through optical fiber in the hand piece to the removable fiber tips. The system may be utilized for a wide variety of surgical and cosmetic procedures.

Indications for Use

The SL3™ is intended to be used for oral soft tissue surgery, including: biopsies, hemostatic assistance; treatment of aphthous ulcers; frenectomy; gingival incision and excision; gingivectomy; gingivoplasty; incising and draining of abscesses; operculectomy; removal of fibromas; soft-tissue crown lengthening; sulcular debridement (removal of diseased or inflamed soft tissue in the periodontal pocket) and tissue retraction for impressions.

What's in the box?

- Laser
- Unifiber™ with Hand Piece
- Foot Pedal
- Disposable Tips (10)
- Power Adapter
- Safety Glasses (3)
- Introductory DVD
- Caution Sign
- Initiating Film
- User's Manual
- Basic User's Guide

What's in the box photo

Section 2: Specifications

LASER SYSTEM SPECIFICATIONS

Size	17.7 cm wide x 13 cm high x 14 cm deep (7" x 5.2" x 5.5")
Weight.....	1.36 kg (3 lbs.)
Wavelength	808 +/- 10 nm
Operation modes	Continuous Wave & Pulsed at 10 Hz
Output power.....	3.0 w ± 20%
Pulse width	50 ms
Calibration	External power meter
Fiber optic port	Standard FT type connector, compatible with 400 µm diameter single-core quartz fiber cable
Aiming beam	650 ± 10 nm red diode laser, 5 mW maximum
Input power	Laser Device: 9V _{DC} , 3.5A / Footswitch: AA Cells, 1.5V
Power supply	100-240 VAC, 1.0 A max, 50-60 Hz, Class 1
Handling/storage.....	Between 0° – 50° C (32° – 122° F) conditions
Protection type	BF part
Footpedal protection	IPX1

Laser System Description

The laser system is contained within a compact lightweight molded plastic housing, consisting of a laser diode assembly with a self-contained cooling system, a flat LCD screen connected to an interface PCB, and power controller PCB.

The base system has two physical buttons. The first button is a mechanical ON-OFF button located on the rear left side of the unit. The second power button is a red LASER STOP button located on the left side. The LASER STOP button de-activates all lasing functions and puts the laser in stand-by mode. A warning message will be displayed. Follow the screen instructions to de-activate the warning message.

The laser output power is regulated in 30 steps by "plus sign" and "minus sign" buttons. The LCD front panel screen indicates the level of output power (in Watts) delivered from the tip of the 400 µm fiber when in continuous wave (CW) mode. An operator can choose CW or Pulsed operation mode by pressing the CW or "Pulse" button on the LCD screen.



The SL3™ uses an internal battery that is recharged when it is plugged in. When fully charged, the battery will last for 30 minutes in continuous use at maximum power. **The SL3™ internal battery is not intended to be changed by the operator.** The laser energy is activated by a wireless foot pedal. Depressing the foot-switch activates the lasing process, while releasing it stops the process. The SL3™ system is highly portable and self-contained with built-in safety features. It is capable of producing the rated output power in a continuous wave mode for more than half an hour without interruption, allowing completion of most surgical procedures. In the pulsed mode, the system delivers pulsed (interrupted) radiation at a repetition rate of 10 Hertz, and a pulse width of 50 ms.

The SL3™ uses disposable, removable fiber-tips to accurately deliver the laser energy. Please refer to page 11 for more information on the disposable tips.

LCD Screen

The SL3™ is equipped with a touch screen, allowing the user to perform selections of various settings. The screen graphics are intuitive, allowing the user to select options quickly and easily. An example of the main touch screen display is shown in the pictures below. For a complete list of all the features on the "Main Display" screen, see page 12.



Unifiber System

Flexible fiberoptic cables are the most widely accepted delivery systems for diode laser power. In the SL3™ system, the 400 µm diameter single-core quartz fiber is mounted into standard FC connectors. The fiber optic cable is coated in a steel metal, offering superior protection and preventing damage to delicate fibers.

The SL3™ uses a Unifiber™ system to deliver diode laser power. This consists of an autoclaveable handpiece shell with internal optics. The hand piece is paired with disposable, non-sterile fiber tips that can be easily removed and replaced in between procedures. For information on setting up the Unifiber system, please refer to page 11.



Section 3: Safety Indications

Testing Standards

The SL3™ Diode Laser has been tested to the standards applicable to medical lasers including IEC60825, IEC60601-1, 60601-2-22, and the Food and Drug Administration's Laser Performance Standard (21 CFR 1040.10 and 1040.11). Please review all labels prior to using the laser.

Familiarity with the American National Standards Institute Z136.3 Document (and/or Laser Institute Safety Manual) is strongly recommended.

Note: The laser must be installed and operated according to CAN/CSA-Z386-92: Laser safety in health care facilities. The labels on the unit are required under these standards for safety purposes and should not be removed.

Labels

Caution Laser in Use

Each treatment area should have a “**Laser in Use**” warning sign posted at the entrance to the treatment area. This signage serves to warn people not to enter the treatment area without proper safety eyewear when the laser is in use.

The Nominal Ocular Hazard Distance with appropriate safety glasses is 1.5 meters or 5 feet.

Class 4 Laser Product.....	808nm ± 10nm 3.0 Watts
Aiming Beam Laser	650nm ± 10nm 5mW max
Pulse Duration.....	50 ms
Beam Divergence.....	617 mRad

Precautions

CAUTION: US Federal law restricts this device to sale by or on the order of a physician.

DO NOT attempt to operate the laser system with any protective panels removed or if the fiber delivery system is improperly connected. The system is equipped with an interlock device for the protective housing/cover.

CAUTION: Do not attempt to remove the cover from the laser chassis for the purpose of repairing the laser. Serious injury from an electrical shock or laser radiation could occur. Removing the cover on the laser chassis will void the warranty.

CAUTION: Not using the proper disposable tip may cause damage to the internal components of the laser.

CAUTION: Changes or modifications not expressly approved by Discus Dental, LLC could void the product warranty.

CAUTION: The remote interlock feature is disabled by default, so the laser is functional. To use this feature, it must be enabled through the “Tools” screen, and additional hardware is required.

CAUTION: Avoid prolonged exposure to the laser energy when working in and around the cervical areas of the tooth. Due to the thin layer of enamel in this area, the laser's energy may be absorbed by the hemoglobin in the pulp in which case pulpal hyperemia may occur. Extended exposure to laser energy could lead to pain and possible pulpal necrosis.

CAUTION: Use a high-volume vacuum to remove the laser “plume.” Provide high-filtration masks for all personnel in the treatment area. The laser plume may contain viable tissue particulates.

CAUTION: The fiber optic cable is very fragile and can be damaged where it attaches to the rear panel. Please be particularly careful when transporting or placing the unit near protruding objects, so as not to damage the cable.

CAUTION: Do not touch the end of the fiber connector or place it on a dusty/dirty surface prior to connecting to the rear panel. If contamination occurs, wipe the connector end with lint free soft tissue dampened with 75-80% ethanol or isopropyl alcohol. Make sure its dry and then attach the connector to the laser port.

CAUTION: This device should be protected against unqualified use when not in use.

DO limit access to the operation area to personnel who are trained in the principles of laser safety. The laser system has a remote interlock option that can be activated if necessary.

DO NOT attempt to defeat the system's interlock or access the enclosures, as they are designed for your protection.

DO NOT attempt repairs of this system. Service and maintenance should only be performed by a qualified SL3™ Service Technician.

CAUTION: Avoid inadvertent laser firing. Turn the laser OFF when not in use for an extended period of time.

Warnings

WARNING: Laser Radiation – Avoid exposure to the eyes or skin from direct or scattered radiation.

DO NOT place any part of the body in direct line with the laser beam.

WARNING: Use of controls or adjustments or performance of procedures other than those specified herein may result in hazardous radiation exposure.

WARNING: Eyewear that protects your eyes from wavelengths other than 808nm do not provide proper protection for use with this laser. Damage to the retina or cornea may be irreparable if exposed to direct, reflected or scattered radiation. Always wear protective eyewear when operating the laser.

WARNING: For the safety of the operator, **DO NOT** use the laser when the fiber is disconnected at the base connection port..

WARNING: Never operate the laser without an attached optical fiber to avoid uncontrolled laser radiation.

WARNING: This laser system is very sensitive to back reflections into the laser module, which may damage or completely destroy the facet of the laser diode. Never point the focused laser beam vertically on a reflecting surface.

WARNING: Laser treatment may result in inadvertent exposure of adjacent tissues. Undue exposure can result in damage to tissue, vessel perforation and bleeding. The practitioner should always set the laser system for minimal exposure to the patient. Optimal parameters for laser surgery may be achieved by starting with the power as low as possible and increasing incrementally as necessary. Power levels affect the precision of cutting, rate of tissue removal and thermal damage to adjacent tissues.

WARNING: The laser can ignite non-metallic materials. All combustible materials must be removed from the operations area or should be kept moist during the procedure.

WARNING: The laser can ignite preparation solutions containing alcohol, acetone or other solvents. **DO NOT** leave puddles of preparation solution in the operations area. Vapors may build up under surgical drapes and create a safety hazard.

DO NOT place the foot-switch in an area where it may be accidentally depressed. When the laser is not in use, remove the foot switch from the practitioner's immediate operations area.

WARNING: Never use the laser system in the presence of flammable anesthetic gases. The use of laser-safe endotracheal tubes and other laser-safe accessories is recommended. Many materials not normally considered flammable could be ignited in the presence of high oxygen and nitrous gas mixtures. An acute awareness of the buildup of the gases as a hazard should be maintained.

WARNING: Avoid tissue splatter on the working end of the disposable tip, as this will create localized heating, which may cause the fiber tip to char and fail. If back splatter occurs, wipe the tip with alcohol gauze. Allow alcohol to evaporate before continuing the lasing process.

DO NOT Use any power supply, other than the one supplied by Discus Dental with your laser. (Part number 04-1473)

Protective Eyewear

All personnel in the operation area, including the patient, must wear eye protection! This is the only patient contra-indication or precaution. Contact lenses are not viable protection. Eye protection must be specific to the wavelength in use (808 +/- 10 nm).

All laser safety glasses/goggles have a wavelength for indicated use stamped into the lens or eyepiece. Care must be taken to assure that everyone in the operations area is wearing the appropriate glasses. (Protective eye wear is marked with optical density >10 at 808 nm).

The laser beam from a laser diode is usually not visible to the human eye, but it can seriously damage retinal tissue. **DO NOT** look directly into the laser beam aperture or into the working end of the optical fiber. Reflected laser light may also cause retinal damage. The reflection hazard exists several feet from the reflection point. Avoid aiming the laser beam in the direction of reflective surfaces.



Laser System Safety Features

The SL3 provides the following safety features for both the user and the patient:

Audible Lasing Signal - An audible signal (high pitch beeping) sounds whenever the activation foot-switch is depressed. (This feature can be adjusted at the Tools section).

Laser Firing Delay - There is a brief delay between depressing the foot switch and the onset of laser activation. This allows the user adequate time to react if the foot switch is inadvertently depressed.

Software Key Lock - The laser system is equipped with a software key-switch to protect against unqualified use, and cannot be turned ON until the correct password is used.

Remote Interlock - The remote interlock feature gives you the option of installing the laser in a dedicated room and be interlocked with the entrance door to the room. In an interlocked installation, the laser shuts off whenever the door is opened, thereby protecting the eyes of the person entering the room. The code to enable the remote interlock is "6 3 2 4". Once the interlock is activated and the proper hardware is connected, the laser will shut off if the remote interlock sensor is triggered. Press the power button to turn the laser back on.

Section 4: System Set Up

Unpacking

Immediately upon receipt of the SL3™ system, inspect all components for damage and for missing items. Unpack all components carefully and verify the presence of all components on the packing slip. Notify Discus Dental immediately if there are any missing components.

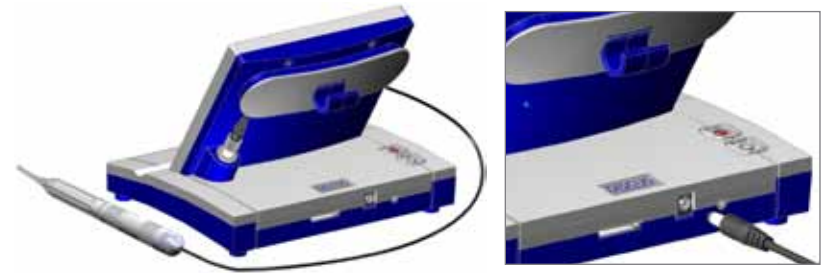
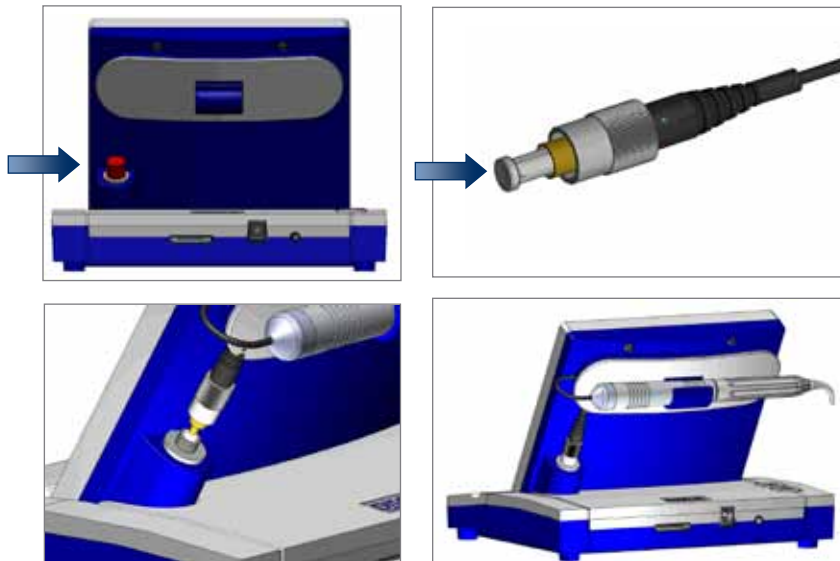
NOTE: Please keep the shipping box during the laser warranty period for possible service, upgrades or returns.

What's in the box?

- Laser
- Unifiber™ with Hand Piece
- Foot Pedal (w/2 AA batteries)
- Disposable Tips (10)
- Power Cable
- Safety Glasses (3)
- Procedural DVD
- Caution Sign
- Initiating Film
- User's Guide
- Basic User's Guide

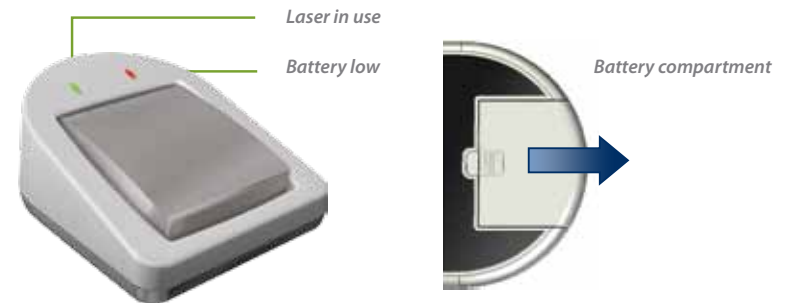
Installation

Place the laser on a suitable table, cart or shelf. Remove the protective end-cap from the laser port (red or white cap). Remove the plastic cap from the fiber FC connector. Attach the Unifiber System into the laser port by turning it clockwise. Wind the fiber cable in a clockwise direction around the spool behind the LCD screen as shown below.



Plug the power cord into a standard AC grounded power outlet (100-240 VAC). Connect the power cord to the proper receptacles on the rear of the laser. Please note that the wide, flat connector to the left of the DC outlet is for factory use only.

The battery-powered wireless foot pedal is composed of a supporting plastic base that is connected to the pedal. In order to power the foot pedal, you must insert the provided AA batteries. Before inserting the batteries, make sure the laser is completely turned off and that the laser is not plugged in.



1. Turn the foot pedal upside down to locate the battery slot
2. Remove the lid and insert the batteries
3. Close the battery lid

The foot pedal has two LED indicator lights located on the top surface of the foot pedal. The green LED indicates that the foot pedal is being depressed and the laser is firing. The amber LED light indicates that the foot pedal battery is low and needs to be replaced.

NOTE: Keep extra AA batteries in your office inventory because the battery should be replaced after approximately 100 hours of continued operation.

NOTE: Some models come with a foot pedal safety shroud (installed separately).

Push the power button on the rear of the laser to the ON position. You will be instructed to input a software lock to protect against unqualified use. Using the keypad on screen, type in "1-2-3-4." You will see a visual verification that the password has been accepted. The panel LCD screen should now read "0.0".



CAUTION: Do not allow the laser system to be exposed to temperatures below 0° C (32° F).

CAUTION: Keep the end of the handpiece clean. Do not allow dust or particles to contaminate the end.

Preparing the Unifiber™ Hand Piece Assembly

The Unifiber™ System is composed of an aluminum autoclavable handle, inner plastic optics and disposable tip. **Please do not under any circumstances autoclave the inner optics sleeve.** Doing so will permanently damage your Unifiber Hand Piece. Connect the inner optics inside the aluminum handle and screw it completely shut. Insert the disposable tip onto the aluminum handle until the magnet sucks the tip in place. Attach the end of the fiber into the fiber port on the back of the SL3's display panel (see Fig. 8). Wrap the fiber around the oval fiber storage plate. There is a rubber ring under the plate to keep the fiber from unraveling. Leave enough fiber unwrapped to reach the patient. The hand piece is now ready for the procedure.

CAUTION : Do not try to detach the fiber from the hand piece, lock-nut or optics.

Cleaning and Sterilization

The SL3™ diode laser system is not supplied in sterile condition, nor must it be sterilized before use. The disposable plastic tips are supplied non-sterile and should be wiped with CaviWipes®, or similar product, prior to use. There is no re-use or re-sterilization procedure indicated. After a used tip is taken off the handpiece, discard it in an infectious waste container (SHARPS).

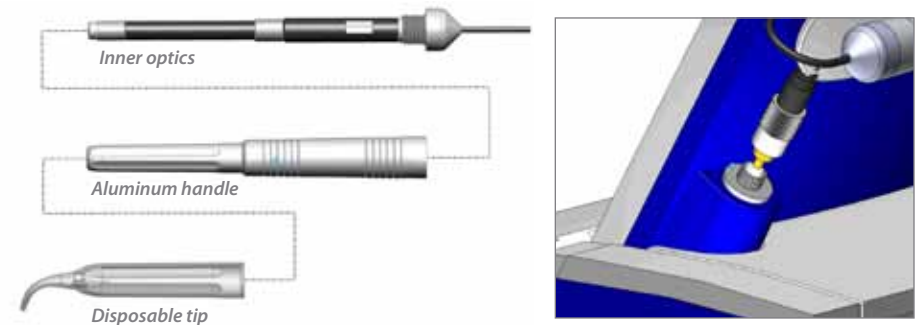
The aluminum hand piece shell is autoclavable and must be sterilized after each use. We recommend a steam sterilization method, performed at 121° C (250° F) at 103.4 kPa (15 PSI) for a minimum of 20 minutes.

NOTE: The LCD screen is not routinely contaminated by the procedure, but the entire front panel should be covered with a protective clear adhesive barrier film, replaceable after each patient. In the event that the screen is touched without protective film, it should be wiped down with CaviWipes® (or equivalent product) and re-covered with a new protective plastic cover.

Transportation

In the event that the laser system is to be relocated (this does not include moving within the facility), place the system into an appropriate shipping carton.

1. Unwind the Unifiber from the rear panel and disconnect it. **Cover all fiber connectors with appropriate end-caps and place the Unifiber into its shipping box.**
2. Disconnect the power supply from the rear of the laser.
3. Carefully repack the laser and attachments into the appropriate shipping cartons.



Section 5: Operation

Procedures Guide

In order to better assist you with your procedures needs, we created a spreadsheet to guide you with the **most conservative** power of the following procedures. The settings shown below are for your guidance only.

Procedure	Technique	Tip	Mode	Power
Abscess	Contact	Initiated	Continuous	1.0 W
Apthous Ulcer	Non-contact	Non-initiated	Pulse	1.4 W
Biopsy	Contact	Initiated	Continuous	1.0 W
Contouring	Contact	Initiated	Continuous	1.0 W
Distal Wedge	Contact	Initiated	Continuous	1.5 W
Draining Abscesses	Contact	Initiated	Continuous	0.9 W
Expose Teeth	Contact	Initiated	Continuous	1.0 W
Fibroma	Contact	Initiated	Continuous	1.0 W
Frenectomy	Contact	Initiated	Continuous	1.1 W
Gingivectomy	Contact	Initiated	Continuous	1.0 W
Gingivitis	Contact	Initiated	Continuous	0.4 W
Gingivitis Therapy	Contact	Initiated	Continuous	0.8 W
Gingivoplasty	Contact	Initiated	Continuous	1.0 W
Hemostasis	Contact	Initiated	Continuous	1.3 W
Herpetic Lesion	Non-contact	Non-initiated	Pulse	1.6 W
Hypertrophic Tissue	Contact	Initiated	Continuous	1.5 W
Implant Exposure	Contact	Initiated	Continuous	1.4 W
Lesion Removal	Contact	Initiated	Continuous	1.0 W
Operculectomy	Contact	Initiated	Continuous	1.5 W
Ovate Pontic	Contact	Initiated	Continuous	1.0 W
Periimplantitis	Contact	Initiated	Continuous	1.0 W
Socket Treatment	Non-contact	Non-initiated	Continuous	0.8 W
Sulcular Debridement	Contact	Initiated	Continuous	0.8 W
Tissue Tag	Contact	Initiated	Continuous	1.5 W
Tissue Welding	Contact	Initiated	Continuous	0.6 W
Tongue Tie	Contact	Initiated	Continuous	1.0 W
Troughing	Contact	Initiated	Continuous	0.8 W
Vestibuloplasty	Contact	Initiated	Continuous	1.0 W

Laser Operation Procedure

It is imperative that you follow steps A through H to ensure that you are protecting yourself, the patient, and staff and to assure that you are using the laser instrument according to the manufacturer's specifications.

- Place appropriate laser-caution signs at all entrances to the operator.
- Verify that the connectors for the power cord (optional) and Unifiber™ at the rear panel are properly secured.
- Prepare appropriate wavelength specific (OD>10 at 808 nm) eyewear. Make sure the operator, assistants and patient are all wearing protective eyewear.
- Turn the laser on by pressing the blue power button located at the back-left panel of the device.
 - Set laser operating parameters manually or use the pre-set procedures.
 - To use pre-set procedures, select the "Presets" button at the bottom of the screen. Navigate to the desired procedure and select it. The laser will set all parameters to the proper settings.
 - If you need to adjust any of the operation settings, click the "Tools" button at the bottom of the screen. This allows you to change the aiming light brightness, beeping volume, voice confirmation volume and the brightness of the screen and remote interlock system.



4. The battery level is displayed on the top right of the screen and demonstrates what percentage of battery life is left. We recommend plugging in the SL3™ if the battery level falls below 25%.
5. Once the correct procedure settings have been selected, the laser is ready to use. Ensure that the proper safety eyewear is worn, and depress the foot pedal to activate the laser energy. Releasing your foot off of the pedal will cause the laser to stop firing.
- E. If the procedure requires an initiated tip, increase the laser output power to its value of 0.6 watts by pressing the “Plus Sign” button. With the tip of the fiber touch a piece of initiating film (dull side is best), and depress the foot switch to activate the laser. An audible high pitch tone will be heard, indicating activation of the laser power. When laser emission begins, move the tip back and forth over the surface, melting the film onto the tip. There will be a rapid melting/ vaporization of the film. The fiber tip should now be dark.
- F. The operator may now proceed with tissue treatment in accordance with developed treatment protocols.
- G. In order to prevent accidental firing, press the Standby/Active button on screen when the laser is not in use. After all sterilizing procedures are completed, press the ACTIVE button and proceed to the next procedure (this laser system does not require warm-up time).
- H. When treatment has been completed, deactivate the laser and push the power button to shut the laser off. Remove the disposable tip and discard it in a Sharps container.



Standby mode

Active

Firing

Section 6: Labels & Symbols

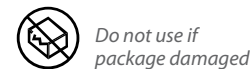
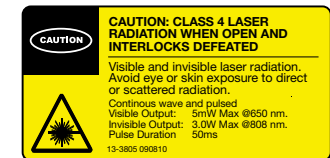
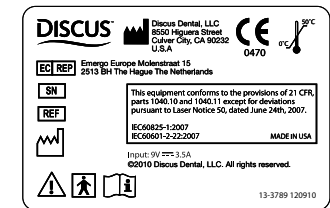
Aperture Label:



Laser Aperture:



Label:



Section 7: Warranty and Service

Service and Maintenance

We suggest that your practice establish an internal verification program for your laser. Verification is recommended a minimum of once per year based on average usage. You may purchase a calibrated hand held power meter approved for use with 808nm devices to check power output. The appropriate safety goggles should be worn at all times when the laser is on. The laser should be set in continuous wave mode. The SL3™ should be set at 0.5, 1.0, 1.5, 2.0 and 2.5 Watts with the output checked at each level. The output display should be within 20% of the meter reading. If not, rescore the fiber and re-check. If the output display is outside the 20% tolerance, return the unit to the manufacturer for recalibration. There are no methods available for the user to adjust the calibration of the unit, and the unit chassis must not be removed by the user for any reason.

SL3™ Diode Laser Unit Warranty

The SL3 Diode Laser Unit (the “Instrument”) manufactured by Discus Dental, LLC (“Discus”) is warranted against defects arising from faulty materials or workmanship for a period of twenty-four (24) months from the date of purchase (the “Warranty Period”). The limited warranties provided herein are expressly conditioned on the Instrument being used under normal conditions and in compliance with the instructions as stated in the Instrument User’s Manual and Basic User’s Guide. The limited warranties provided herein do not cover any damage that may occur as a result of misuse, neglect, adjustments, or alterations to the Instrument.

The warranty offered by Discus on the Instrument is a limited warranty and the sole liability of Discus shall be to replace or repair the Instrument. Discus shall have no obligation or liability to refund any portion of the purchase price and shall have no liability for special, exemplary, consequential or punitive damages, loss of profits, damages to persons or injury in connection with the purchase or use of the Instrument. The Warranty does not include labor, postage, or delivery charges. This warranty does not apply to the external finish of the console, handpiece, fiber, power cord, or foot pedal. Discus reserves the right to make changes in design or to modify such previously manufactured products, at its sole discretion.

Use of products not specifically made by or authorized by Discus for use with the Instrument will void and nullify all warranties. The Instrument is not to be serviced without specific written instructions from Discus. Any attempted service of the Instrument without these instructions will void the Discus Warranty.

Warranty Evaluation & Repair Service

Subject to the determination set forth below, if a repair is required during the Warranty Period, Discus shall authorize warranty repair or replacement at no cost to you. Please contact our Customer Service Department TOLL FREE

at: 1.800.217.8822 to request warranty repair service and obtain a Return Merchandise Authorization (RMA) number for shipping purposes. Please ensure that the RMA number is clearly marked on the box used to return the Instrument. It is recommended that you return your Instrument in its original shipping box. If the original shipping box is not available, make sure that adequate packing is used to protect against shipping damage. Make sure you buy insurance to protect your asset against shipping damage. Please clearly state the reasons for return.

Note: Returns without a return authorization number will be refused.

Send Returns to: Attention Shipping Manager
Discus Dental, LLC Laser Repair Department
1700 S. Baker Ave.
Ontario, CA 91761
(888) 876-4546

Discus will inspect your Instrument and determine whether the damage and/or failure are covered under the Warranty. Warranty repairs or replacement will be completed and the Instrument will be shipped back to you within five (5) business days of receipt of the Instrument.

If the required repairs are not covered by the Warranty due to, in Discus’s determination, misuse, neglect, adjustments or alterations to the Instrument, or the applicable Warranty Period having ended, you will be contacted within two (2) business days of receipt of the Instrument with a price quotation for the cost of repair(s) or cost of the replacement. Upon receipt of your approval and acceptance of repair and shipping costs, Discus will repair and ship your Instrument back to you within five (5) business days.

Replacement/Repair Program & Services

Discus provides for the purchase of major Instrument components after the warranty periods have ended. We guarantee the availability of replacement parts for a period of one year after sale of this product is discontinued, should that occur.

Refurbished parts, if available, are offered to our customers at a lower cost. Refurbished parts are those that have been returned to us, remanufactured and inspected to ensure they meet the original performance standards for new components before being offered for resale. Discus provides a new warranty on all new or refurbished parts purchased. The warranty periods for the newly purchased parts begin when the new or refurbished part is shipped.

Non -Warranty Evaluation & Repair Service

Please contact our Customer Service Department to request repair service outside the Warranty Period. Our Customer Service Department will provide a RMA. The customer is responsible for freight charges to ship the Instrument to and from Discus for non-warranty service. Returns without a RMA will be refused.

A minimum Evaluation Fee will be charged for all repairs not covered by warranty. If the customer does not authorize the repair, the customer is still responsible for payment of the minimum Evaluation Fee. If the customer authorizes repair service, the minimum Evaluation Fee will be applied to the total repair costs.

Upon receipt of approval and acceptance of the repair and shipping costs, Discus will repair and ship the Instrument back within five (5) business days.

Terms & Conditions

1. All prices subject to change without notice.
2. All prices subject to applicable sales tax.
3. Refurbished units subject to availability.
4. Returning parts after warranty period may be subject to an evaluation fee.
5. All program information is based on the latest data available and is subject to change.

For questions regarding defective product during the warranty period, please call Discus Customer Service or an authorized sales distributor.

Address: Discus Dental, LLC	USA Toll Free: (800) 422-9448
8550 Higuera Street	Canada Toll Free: (800) 278-8282
Culver City, CA 90232	For all other countries, please see User's Manual
Tel: (310) 845-8200	or visit: www.discusdental.com

Compliance

FCC Statement

This device complies with Part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

This product has been tested and complies with the specifications for a Class B digital device, pursuant to Part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates, uses, and can radiate radio frequency energy and, if not installed and used according to the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to radio or television reception, which is found by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:

- Reorient or relocate the receiving antenna
- Increase the separation between the equipment or devices
- Connect the equipment to an outlet other than the receiver's
- Consult a dealer or an experienced RF technician for assistance

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

Industry Canada Statement

This device complies with Industry Canada license-exempt RSS standard(s). Operation is subject to the following two conditions: (1) this device may not cause interference, and (2) this device must accept any interference, including interference that may cause undesired operation of the device.

Le présent appareil est conforme aux CNR d'Industrie Canada applicables aux appareils radio exempts de licence. L'exploitation est autorisée aux deux conditions suivantes : (1) l'appareil ne doit pas produire de brouillage, et (2) l'utilisateur de l'appareil doit accepter tout brouillage radioélectrique subi, même si le brouillage est susceptible d'en compromettre le fonctionnement.

If you have questions or comments about the contents of this User's Manual, please call our toll free number for assistance: (800) 422-9448. We want to make sure your experience in using our SL3™ system is most gratifying.

Notes
