

**510(k) Summary of Safety and Effectiveness
Sharplan Lasers, Inc. 15F CO2 Laser System
an Expanded Indication for Use**

In response to the Safe Medical Devices Act of 1990, the following is a summary of the safety and effectiveness information upon which the substantial equivalence determination is based.

The safety and effectiveness of Sharplan 15F CO2 Laser System for the intended use as a heat to activate tooth bleaching materials is based upon a determination of the substantial equivalence as well as the safety and effectiveness of its predicate devices which include the Sharplan 15F CO2 Laser and the Ion Laser Technology (ILT) CO2 Laser System.

1. Sharplan Lasers, Inc.
1 Pearl Court
Allendale, NJ 07401
George J. Hattub, Director of Regulatory Affairs/Quality Assurance
April 30, 1997
2. **Model:** Sharplan Model 15F CO2 Laser
3. **Predicate Devices:** The Sharplan 15 F CO2 Laser (K950725) and the ILT CO2 Laser System (K954460).
4. **Description:** The Sharplan Model 15F CO2 Surgical Laser is a medical device which capable of being used as a heat source for activating tooth bleaching materials. It emits a treatment laser beam at a wavelength of 10.6 μm up to 15 watts in either continuous or pulse wave modes of operation.
5. The Sharplan Model 15F (for this indication) is substantially equivalent to the following devices: The Sharplan 15F CO2 Laser which is intended for use in ENT and soft tissue dentistry and the ILT Surgical Laser which is intended for use as a heat source to activate tooth bleaching materials, as well as soft tissue incision and ablation in ENT, and soft tissue dentistry.

No new intended use for a CO2 Laser was sought in this premarket notification. Therefore, no clinical data was presented.

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(continued)**

- 6 . From a design and clinical perspective, the predicate and candidate laser devices, are of the same technology (one device is identical), and have the same intended use. Based upon an analysis of the overall performance characteristics for the devices, Sharplan Lasers, Inc. believes that no significant differences exist. Furthermore, this expanded indication does not raise any concerns regarding the safety or effectiveness of the Sharplan Lasers, Inc. 15F CO₂ Laser System for use as a heat source to activate tooth bleaching materials.

Advisory: This information was prepared for the sole purpose of compliance with the Safe Medical Devices Act of 1990. It does not imply that the procedures described herein can be performed with the equipment described without substantial risk of personal injury or death to patients due to operator error or in procedures requiring a high degree of skill.



Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Mr. George J. Hattub
Director Regulatory Affairs/Quality Assurance, CQE
Sharplan Lasers, Inc.
1 Pearl Court
Allendale, New Jersey 07401

JUL 18 1997

Re: K971743
Trade Name: Sharplan Model 15F CO₂ Laser
Regulatory Class: II
Product Code: GEX
Dated: May 9, 1997
Received: May 12, 1997

Dear Mr. Hattub:

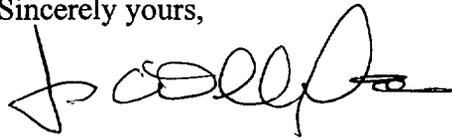
We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic (QS) inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4595. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

Sincerely yours,



f Celia M. Witten, Ph.D., M.D.
Director
Division of General and
Restorative Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number (If known): K971743

Device Name: Sharplan 15F CO2 Laser System

Indications For Use:

The Sharplan 15F CO2 Laser System is intended for use as a heat source to activate tooth bleaching materials.

(Please Do Not Write Below This Line - Continue on Another Page If Needed)

Concurrence of CDRH, Office of Device Evaluation (ODE)



(Division Sign-Off)

Division of General Restorative Devices

510(k) Number

K971743

Prescription Use
(Per 21 CFR 801.109)

OR

Over-The-Counter Use

(Optional Format 1-2-96)