K072424

OCT 3 2007

Premarket Notification 510(k) Summary
CoolTouch LC215 Nd:YAG Laser System

This 510(K) Summary of safety and effectiveness for the CoolTouch LC215 Nd:YAG surgical laser system is submitted in accordance with the requirements of 21CFR 807.92.

Applicant:	New Star Lasers, Inc. dba CoolTouch, Inc.
Address:	9085 Foothills Boulevard Roseville, CA 95747
Contact Person:	Natalie R. Vollrath
Telephone:	(916) 677-1900
Fax:	(916) 677-1901
Preparation Date:	August 27, 2007
Device Trade Name:	CoolTouch LC215 Nd:YAG Surgical Laser
Common Name:	Nd:YAG Pulsed Surgical Laser
Classification Name:	Instrument, Surgical Powered, Laser 79-GEX
Legally Marketed Predicate Device:	CoolTouch CT3S Nd:YAG Laser System
Description of the CoolTouch LC215 Nd:YAG Surgical Laser:	The LC215 Nd:YAG Laser System produces laser emission at 1320nm. The laser consists of three interconnected sections: the cabinet which houses the power supply, the cooling system, the microcontroller and the laser, the fiber optics, and the handpiece or JouleTracker
Intended use of the CoolTouch LC215 Nd:YAG Surgical Laser:	For use in dermatology for incision, excision, ablation and vaporization with hemostasis of soft tissue. For use in the treatment of fine lines and wrinkles. For treatment of back acne and atrophic acne scars. For treatment of reflux of the great and small saphenous veins associated with varicose veins and varicosities
Performance Data:	None
Conclusion:	Based on the evaluation of the risks and hazards and including various testing of the modifications, the CoolTouch LC215 Nd:YAG Surgical Laser System is substantially equivalent to the predicate device, the CT3S

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

New Star Lasers, Inc. dba
CoolTouch, Inc.
% Ms. Natalie Vollrath
Quality Assurance Manager
9085 Foothills Boulevard
Roseville, California 95747

OCT 3 ˇ 2007

Re: K072424
Trade/Device Name: CoolTouch LC215
Regulation Number: 21 CFR 878.4810
Regulation Name: Laser surgical instrument for use in general and plastic surgery and in dermatology
Regulatory Class: II
Product Code: GEX
Dated: September 18, 2007
Received: September 19, 2007

Dear Ms. Vollrath:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate 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) that do not require approval of a premarket approval application (PMA). 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 (PMA), 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 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the

This letter will allow you to begin marketing your device as described in your Section 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), please contact the Office of Compliance at (240) 276-0115. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,

Mark N. Melkerson
Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number Pending

Device Name CoolTouch LC215

Indications for Use

The CoolTouch Model LC215 Nd:YAG Surgical Laser is indicated for the following:

a) for use in dermatology for incision, excision, ablation and vaporization with hemostasis of soft tissue;

b) for use in the treatment of fine lines and wrinkles;

c) for treatment of back acne and atrophic acne scars, and;

d) for treatment of reflux of the great and small saphenous veins associated with varicose veins and varicosities.

(Division Sign-Off)
Division of General, Restorative, and Neurological Devices

510(k) Number K072424

Prescription Use ___X___ AND/OR Over-the-Counter Use _____
(Part 21 CFR 801 Subpart D) (21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)