

Attachment I
510(K) Summary
ProLite Pulsed Light System

JAN 8 2002

K013365

This 510(K) Summary of safety and effectiveness for the ProLite VL Pulsed Light System is submitted in accordance with the requirements of the SMDA 1990 and following guidance concerning the organization and content of a 510(K) summary.

Applicant: Medical Bio Care Sweden AB.

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Preparation Date: October 5, 2001

Device Trade Name: ProLite Pulsed Light System

Common Name: Intense Pulsed Light System

Classification Name: Instrument, Surgical, Powered, laser
79-GEX, 21 CFR 878-48

Legally Marketed Predicate Device: Photoderm PL System
K number K60772

Description of the ProLite V Pulsed Light System The ProLite Pulsed Light System delivers pulsed light at a wavelength of 550 nanometers. The device consists of three interconnected sections: The cabinet which houses the power supply, the cooling system and the microcontroller, the umbilical to the handpiece, and the handpiece, which houses the waveguide

Intended use of the ProLite V Pulsed Light System The ProLite Pulsed Light System is indicated the treatment of benign pigmented lesions and the removal of tattoos.

Performance Data: None

Conclusion: The ProLite Pulsed Light System is substantially equivalent to other existing pulsed light systems in commercial distribution for treatment of benign pigmented lesions and the removal of tattoos in Dermatology and Plastic Surgery.



JAN 8 2002

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Medical BioCare Sweden AB
c/o Ms. Connie White Hoy
908 Stetson Street
Woodland, California 95776

Re: K013365

Trade/Device Name: ProLite Pulsed Light System

Regulation Number: 878.4810

Regulation Name: Laser surgical instrument for use in general
and plastic surgery and in dermatology

Regulatory Class: II

Product Code: GEX

Dated: October 1, 2001

Received: October 10, 2001

Dear Ms. Hoy:

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.

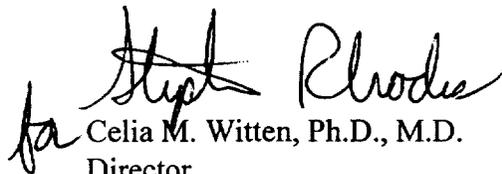
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's 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 quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

Page 2 – Ms. Connie White Hoy

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 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4659. 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" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>

Sincerely yours,

Handwritten signature of Celia M. Witten in black ink, appearing as 'Celia M. Witten'.

Celia M. Witten, Ph.D., M.D.

Director

Division of General, Restorative
and Neurological Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

Enclosure

K013365/A1

INDICATION FOR USE STATEMENT

510(k) Number: K013365

Device Name: ProLite Pulsed Light System

Indications for Use:

The ProLite Pulsed Light System is intended for the treatment of benign pigmented lesions and the removal of tattoos.

(Please do not write below this line - Continue on another page if needed)

Concurrence of CDREH, Office of Device Evaluation (ODE)

[Handwritten Signature]

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

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Nov 3 10 15 AM '01

FDA

510(k) Number K013365

Prescription Use
(per 21 CFR 801.109)

OR

Over-the-Counter Use