



Food and Drug Administration
10903 New Hampshire Avenue
Document Control Center – WO66-G609
Silver Spring, MD 20993-0002

LED Intellectual Properties LLC
Mr. Steve Marchese
Chief Executive Officer
16552 Von Karman Avenue
Irvine, California 92606

October 23, 2015

Re: K151336
Trade/Device Name: LightStim Professional 2-Panel Light
Regulation Number: 21 CFR 878.4810
Regulation Name: Laser surgical instrument for use in general and plastic surgery and in dermatology
Regulatory Class: Class II
Product Code: OHS, OLP
Dated: May 5, 2015
Received: September 16, 2015

Dear Mr. Marchese:

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. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you; however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to 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); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); 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.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to

<http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

Joshua C. Nipper -S

For Binita S. Ashar, M.D., M.B.A., F.A.C.S.
Director
Division of Surgical Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indications for Use

510(k) Number (if known)

K151336

Device Name

LightStim Professional 2-Panel Light

Indications for Use (Describe)

The LightStim Professional System #3A panel is intended for the use in the treatment of full-face wrinkles.

The LightStim Professional System #3B panel is intended for the use in the treatment of mild to moderate acne.

The LightStim Professional System #4 panel is intended for the use in the treatment of mild to moderate acne.

Type of Use (Select one or both, as applicable)

Prescription Use (Part 21 CFR 801 Subpart D)

Over-The-Counter Use (21 CFR 801 Subpart C)

CONTINUE ON A SEPARATE PAGE IF NEEDED.

This section applies only to requirements of the Paperwork Reduction Act of 1995.

DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.

The burden time for this collection of information is estimated to average 79 hours per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to:

Department of Health and Human Services
Food and Drug Administration
Office of Chief Information Officer
Paperwork Reduction Act (PRA) Staff
PRASStaff@fda.hhs.gov

"An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number."

510(k) Summary of Safety and Effectiveness

LED Intellectual Properties, LLC.

Device: LightStim Professional 2-Panel Light System

1. General Information

Submitter: LED Intellectual Properties, LLC
16552 Von Karman Ave. Irvine, Ca. 92606

Contact Person: Steve Marchese & Chase Marchese
Office: (949) 502-4088 Mobile: (714) 924-0492

Emails: Steve@lightstim.com & Chase@lightstim.com

2. Device name and code

Device Proprietary Name: LightStim Professional 2-Panel Light System

Classification Code and name: OHS – Light Based OTC Wrinkles and OLP – OTC Light Based For Acne

3. Predicate Devices

LED Intellectual Properties LLC LightStim Professional 2-Panel Light (K150098), LightStim for Acne + LightStim for Acne Mini (K142246), LightStim For Acne (K131461), LightStim for Wrinkles (K120775)

4. Device Description

The LightStim Professional 2-Panel Light System is a portable device that has two (2) separate interchangeable 2-panel treatment heads, each containing 1,130 LEDs. Each treatment heads is attached to a fully retractable and articulating arm with a 42" reach. Either one of these arms can be attached to the rolling stand for use. There is an on/off switch, and an AC to DC power supply.

The LightStim 2-Panel Light System is intended to be operated only by a person who has been personally trained by Company personnel. Each device includes a detailed Instruction Manual and includes one-on-one training and instruction by Company training staff. That training includes a thorough review of the Instruction Manual for the device and actual one-on-one training on the operation of the device with the end user.

5. Indications for Use

The LightStim Professional System #3A panel is intended for the use in the treatment of full-face wrinkles

The LightStim Professional System #3B panel is intended for the use in the treatment of mild to moderate acne.

The LightStim Professional System #4 panel is intended for the use in the treatment of mild to moderate acne.

6. Substantial Equivalency

The LightStim Professional 2- Panel Light System emits visible and/or infrared energy and the intended use is the same as the intended use of the predicate devices. Further, the predicate devices and the LightStim Professional 2-Panel Light System are produced by the same LED Intellectual Properties, LLC, and they utilize LEDs with the similar outputs and technical characteristics

7. Performance Testing

Our device LightStim Professional 2-Panel results in patient contact with an electrically powered component therefore; it was tested for conformance to IEC / EN 60601-1 3rd Edition: Medical Electrical Equipment - Part 1: General requirements for basic safety and essential performance

Our device LightStim Professional 2-Panel includes an electronic component, was tested for conformance to IEC / EN 60601-1-2 3rd Edition Medical Electrical Equipment – Part 1-2: General Requirements for Safety and Essential Performance - Collateral standard: Electromagnetic compatibility – Requirements and tests.

Therefore, taking into consideration the Table for Substantial Equivalence, an analysis of safety, indications, intended uses, performance, and technological properties, the LightStim Professional 2-Panel Light System raises no new issues of safety or effectiveness and has been found to be substantially equivalent to the predicate devices.