



February 28, 2023

Shenzhen Ulike Smart Electronics Co.,Ltd  
% Riley Chen  
Registration Engineer  
Feiyang Drug & Medical Consulting Technical Service Group  
Rm 2401 Zhenye International Business Center, No. 3101-90,  
Qianhai Road  
Shenzhen, 518052  
China

Re: K223618

Trade/Device Name: IPL Hair Removal Device, Model(s): UI06 PN, UI06 PL, UI06 JL, UI06 BR,  
UI06 DB, UI06 PR, UI06 OG, UI06 RD

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: OHT

Dated: December 2, 2022

Received: December 5, 2022

Dear Riley Chen:

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. Although this letter refers to your product as a device, please be aware that some cleared products may instead be combination products. The 510(k) Premarket Notification Database located at <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpmn/pmn.cfm> identifies combination product submissions. 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) for devices or postmarketing safety reporting (21 CFR 4, Subpart B) for combination products (see <https://www.fda.gov/combination-products/guidance-regulatory-information/postmarketing-safety-reporting-combination-products>); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820) for devices or current good manufacturing practices (21 CFR 4, Subpart A) for combination products; and, if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <https://www.fda.gov/medical-devices/medical-device-safety/medical-device-reporting-mdr-how-report-medical-device-problems>.

For comprehensive regulatory information about medical devices and radiation-emitting products, including information about labeling regulations, please see Device Advice (<https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance>) and CDRH Learn (<https://www.fda.gov/training-and-continuing-education/cdrh-learn>). Additionally, you may contact the Division of Industry and Consumer Education (DICE) to ask a question about a specific regulatory topic. See the DICE website (<https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance/contact-us-division-industry-and-consumer-education-dice>) for more information or contact DICE by email ([DICE@fda.hhs.gov](mailto:DICE@fda.hhs.gov)) or phone (1-800-638-2041 or 301-796-7100).

Sincerely,

  
Colin K. Chen -S

for

Jianting Wang  
Acting Assistant Director  
DHT4A: Division of General Surgery Devices  
OHT4: Office of Surgical  
and Infection Control Devices  
Office of Product Evaluation and Quality  
Center for Devices and Radiological Health

Enclosure

## Indications for Use

510(k) Number (if known)  
K223618

Device Name

IPL Hair Removal Device

Model(s): UI06 PL, UI06 PN, UI06 JL, UI06 BR, UI06 DB, UI06 PR, UI06 OG, UI06 RD

Indications for Use (Describe)

IPL Hair Removal Device is indicated for the removal of unwanted hair. The device is also indicated for the permanent reduction in hair regrowth, defined as the long-term, stable reduction in the number of hairs regrowing when measured at 6, 9 and 12 months after the completion of a treatment regime.

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.

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## 510(k) Summary

"510(k) Summary" as required by 21 CFR Part 807.92.

### I. Submitter

Shenzhen Ulike Smart Electronics Co.,Ltd.  
No.8, Keyuan Road, Yuehai Sub-district, Nanshan District, Shenzhen, Guangdong, China  
Post code: 518000  
Tel.: +86 18600825411

Shane Xie  
Registration Director  
Tel: +86 18600825411  
Email: Shane@ulikebeauty.com

### II. Device

Name of Device: IPL Hair Removal Device  
Model(s): UI06 PL, UI06 PN, UI06 JL, UI06 BR, UI06 DB, UI06 PR, UI06 OG, UI06 RD  
Common or Usual Name: Light Based Over-The-Counter Hair Removal  
Regulation Name: Laser surgical instrument for use in general and plastic surgery and in dermatology  
Regulatory Class: II  
Product Code: OHT  
Regulation Number: 21 CFR 878.4810

### III. Predicate Device

Primary predicate device:

<u>Manufacturer</u>	<u>Predicate Device</u>	<u>510(k) Number</u>	<u>Approval Date</u>
CyDen Limited.	iPulse SmoothSkin Gold Hair Removal System	K160968	Apr.14, 2016

Predicate device:

<u>Manufacturer</u>	<u>Predicate Device</u>	<u>510(k) Number</u>	<u>Approval Date</u>
Shen Zhen CosBeauty Co., Ltd	IPL Hair Removal Device Joy Version, CB-027	K173813	Sept. 07, 2018

Reference device:

<u>Manufacturer</u>	<u>Reference Device</u>	<u>510(k) Number</u>	<u>Approval Date</u>
Dongguan Define Beauty Electronic Technology Co. Ltd	IPL HAIR REMOVAL SG-8025	K212318	Jan.14, 2022

#### IV. Device Description

IPL Hair Removal Device (Model: UI06 PL, UI06 PN, UI06 JL, UI06 BR, UI06 DB, UI06 PR, UI06 OG, UI06 RD), is an over-the-counter, home-use and personal device for hair reduction by using Intense Pulsed Light (IPL). It works below the skin's surface and does not involve any cutting or pulling, reducing hair growth with minimal pain.

The device is only powered by the external power adapter and its IPL emission activation is by finger switch. This product adopts irreplaceable sapphire treatment window and the spot size is 3.3cm<sup>2</sup> that is suitable for multiple hair removal areas. It contains a skin sensor to detect appropriate skin contact, if the device is not in full contact with the skin, the device cannot emit the treatment light pulses. Besides, the device has the cooling function that will be activated throughout the whole hair removal process to provide users with a more comfortable experience.

IPL Hair Removal Device, model: UI06 PL, UI06 PN, UI06 JL, UI06 BR, UI06 DB, UI06 PR, UI06 OG, UI06 RD have the same indication for use, performance, structure design and operation, the only difference is their enclosure color (UI06 PL is Purple, UI06 PN is Pink, UI06 JL is Yellow, UI06 BR is Brown, UI06 DB is Dark Brown, UI06 PR is Dark Purple, UI06 OG is Maple Red, UI06 RD is Red).

#### V. Indications for Use

IPL Hair Removal Device is indicated for the removal of unwanted hair. The device is also indicated for the permanent reduction in hair regrowth, defined as the long-term, stable reduction in the number of hairs regrowing when measured at 6, 9 and 12 months after the completion of a treatment regime.

#### VI. Materials

Component name	Material of Component	Body Contact Category	Contact Duration
IPL Hair Removal Device (Enclosure and treatment window)	ABS, PC, Crystal application	Surface-contacting device: Intact skin	Less than 24 hours

We have tested the device for biocompatibility by a reliable third-party lab. For details, please refer to “Biocompatibility Discussion”.

#### VII. Comparison of Technological Characteristics With the Predicate Device

The IPL Hair Removal Device has the same intended use, mode of action and similar operational characteristics as the predicate devices and reference device. Any minor differences between the subject device and the listed predicate devices and reference device do not raise any issues of safety or efficacy. Performance data supports that the device is safe and as effective as the

predicate devices and reference device for its intended use. Therefore, the IPL Hair Removal Device may be found substantially equivalent to its predicate devices and reference device. IPL Hair Removal Device is compared with the following Predicate Devices and Reference Device in terms of intended use, design, specifications and performance:

<u>Comparison Elements</u>	<u>Subject Device</u>	<u>Primary Predicate Device</u>	<u>Predicate Device</u>	<u>Reference Device</u>	<u>Remark</u>
510(k) Number	K223618	K160968	K173813	K212318	/
Trade name	IPL Hair Removal Device	iPulse SmoothSkin Gold Hair Removal System	IPL Hair Removal Device Joy Version, CB-027	IPL HAIR REMOVAL SG-8025	/
Manufacturer	Shenzhen Ulike Smart Electronics Co.,Ltd.	Cyden Limited.	Shen Zhen CosBeauty Co., Ltd	Dongguan Define Beauty Electronic Technology Co. Ltd	/
Regulation number	21 CFR 878.4810	21 CFR 878.4810	21 CFR 878.4810	21 CFR 878.4810	Same
Product code	OHT	OHT, GEX	OHT	OHT	Same
Device classification	Class II	Class II	Class II	Class II	Same
Indication for use/ Intended use	IPL Hair Removal Device is indicated for the removal of unwanted hair. The device is also indicated for the permanent reduction in hair regrowth, defined as the long-term, stable reduction in the number of hairs regrowing when measured at 6, 9 and 12 months after the completion of a treatment regime.	The iPulse SmoothSkin Gold Hair Removal System is indicated for the removal of unwanted hair. The iPulse Smoothskin Gold is also indicated for the permanent reduction in hair regrowth, defined as the long-term, stable reduction in the number of hairs regrowing when measured at 6, 9 and 12 months after the completion of a treatment regime.	The IPL Hair Removal Device Joy Version is indicated for the removal of unwanted hair. The device is also indicated for the permanent reduction in hair regrowth, defined as the long-term, stable reduction in the number of hairs regrowing when measured at 6, 9 and 12 months after the completion of a treatment regime. The device is used for adults with Fitzpatrick skin types I - IV.	The IPL Hair Removal (Model: SG-8025) is an over the Counter device intended for the removal of unwanted body and/or facial hair in adults. It is also intended for permanent reduction in unwanted hair. Permanent hair reduction is defined as the long-term stable reduction in the number of hairs regrowing when measured at 6. 9. and 12 months after the completion of a	Same, only wording difference

<u>Comparison Elements</u>	<u>Subject Device</u>	<u>Primary Predicate Device</u>	<u>Predicate Device</u>	<u>Reference Device</u>	<u>Remark</u>	
				treatment regimen.		
Prescription or OTC	OTC	OTC	OTC	OTC	Same	
Applicable skin	Fitzpatrick Skin Types I-V	Unknown	Fitzpatrick skin types I - IV	Unknown	Different	
Treatment area	large areas (e.g. arms, legs, chest) and small areas (e.g. lip)	Unknown	Large areas (legs, arms, back and abdomen), face (upper lip, chin and sideburns)	Body and face	Similar	
<b>Device design</b>						
Source energy	Supplied by external adapter	External supply	Power	Supplied by external adapter	Supplied by external adapter	Same
Power supply	100~240V, 50/60Hz	110V or 230V, 50/60Hz		100-240 VAC, 50/60Hz	Unknown	Same
Dimension	58×34×179mm (W x H x D)	Unknown		126*78*200mm	205*76*56mm (H*W*D)	Different
Sterilization	Not required	Not required		Not required	Not required	Same
<b>Output specification</b>						
Light source	Intense Pulsed Light	Intense Pulsed Light		Intense Pulsed Light	Intense Pulsed Light	Same
Energy medium	Xenon Arc Flashlamp	Xenon Arc Flashlamp		Xenon Arc Flashlamp	Xenon lamp	Same
Wavelength range	560-1200nm	510-1100nm		510-1200nm	530nm	Similar
Energy density	3~6 J/cm <sup>2</sup>	3-6J/cm <sup>2</sup>		1.8~5.1J/cm <sup>2</sup>	2.5J/cm <sup>2</sup> Max	Similar
Output energy	9.9~19.8J	9~18J		Body lamp cartridge 11.77~22.21J 510-1200nm Facial lamp cartridge 3.65~7.04J 512-1197nm Bikini lamp cartridge 3.84~7.22J 511-1200nm	Level 1: 7.5J Level 2: 8.5J Level 3: 9.5J Level 4: 11J Level 5: 12J	Similar
Spot size	3.3cm <sup>2</sup>	3cm <sup>2</sup> (3cm by 1cm)		Body: 4.2cm <sup>2</sup> Bikini and face: 2.0cm <sup>2</sup>	3.0cm <sup>2</sup>	Similar
Pulse duration	1ms~7ms	2ms to 10ms		9.2~11.2ms	1ms	Similar
Pulsing control	Finger switch	Finger switch		Finger switch	Finger switch	Same

<u>Comparison Elements</u>	<u>Subject Device</u>	<u>Primary Predicate Device</u>	<u>Predicate Device</u>	<u>Reference Device</u>	<u>Remark</u>
Delivery device	Direct illumination to tissue	Direct illumination to tissue	Direct illumination to tissue	Direct illumination to tissue	Same
Output intensity level	3	Unknown	5	5	Different
Software/ Firmware/ Microprocess or Control?	Yes	Yes	Yes	Yes	Same
<b>Additional features</b>					
Electrical safety	ANSI AAMI ES60601-1 IEC 60601-1-2 IEC 60601-1-11 IEC 60601-2-57 IEC 60601-2-83	IEC 60601-1 IEC 60601-1-2 IEC 60601-2-57	IEC 60601-1 IEC 60601-1-2 IEC 60601-2-57	IEC 60601-1 IEC 60601-1-2 IEC 60601-1-11 IEC 60601-2-57	Similar
Eye safety	IEC 62471	IEC 62471	Unknown	IEC 62471	Same
Biocompatibility	ISO 10993-5 ISO 10993-10	ISO 10993-5 ISO 10993-10	ISO 10993-5 ISO 10993-10	ISO 10993-5 ISO 10993-10	Same

### VIII. Performance Data

The following performance data were provided in support of the substantial equivalence determination.

#### **1) Biocompatibility Testing**

The biocompatibility evaluation for the body-contacting components of the IPL Hair Removal Device was conducted in accordance with the “Use of International Standard ISO 10993-1, 'Biological Evaluation of Medical Devices –Part 1: Evaluation and Testing Within a Risk Management Process, Document Issued on September 4, 2020”, as recognized by FDA. The following testing was performed to, and passed, including:

- ISO 10993-5:2009, Biological Evaluation of Medical Devices –Part 5: Tests for In Vitro Cytotoxicity
- ISO 10993-10:2010, Biological Evaluation of Medical Devices –Part 10: Tests for Irritation and Skin Sensitization

#### **2) Electrical Safety and EMC**

Electrical safety and EMC testing was performed to, and passed, as per the following standards:

- ANSI AAMI ES60601-1 Medical electrical equipment - Part 1: General requirements for basic safety and essential performance



- IEC 60601-1-2 Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance - Collateral Standard: Electromagnetic disturbances - Requirements and tests
- IEC 60601-1-11 Medical Electrical Equipment –Part 1-11: General Requirements for Basic Safety and Essential Performance –Collateral Standard: Requirements for Medical Electrical Equipment and Medical Electrical Systems Used in the Home Healthcare Environment
- IEC 60601-2-57 Medical electrical equipment –Part 2-57: Particular requirements for the basic safety and essential performance of non-laser source equipment intended for therapeutic, diagnostic, monitoring and cosmetic/aesthetic use
- IEC 60601-2-83 Medical electrical equipment - Part 2-83: Particular requirements for the basic safety and essential performance of home light therapy equipment

### **3) Eye Safety**

- IEC 62471 Photobiological safety of lamps and lamp systems

### **4) Software Verification and Validation**

Software documentation consistent with *moderate level* of concern was submitted in this 510(k). System validation testing presented in this 510(k) demonstrated that all software requirement specifications are met and all software hazards have been mitigated to acceptable risk levels.

### **5) Usability**

The product usability has been evaluated and verified according to the following standard and FDA guidance

- IEC 60601-1-6 Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability
- Applying Human Factors and Usability Engineering to Medical Devices, issued on FEBRUARY 2016

### **Summary**

Based on the above performance as documented in this application, IPL Hair Removal Device was found to have a safety and effectiveness profile that is similar to the predicate devices and reference device.

## **IX. Conclusions**

Based on the above analysis and non-clinical tests performed, it can be concluded that the subject device IPL Hair Removal Device is as safe, as effective, and performs as well as the legally marketed predicate devices and reference device.