CLEARED

May 20, 2026
86 days to decision

K260594 IPL Hair Removal Device (FL-B505AG, FL-B505AP, FL-B505AM, FL-B507AG, FL-B507AP, FL-B507AM, FL-B508AG, FL-B508AP, FL-B508AM, FL-B509AG, FL-B509AP, FL-B509AM, FL-B510G, FL-B510P, FL-B510M, FL-B511G, FL-B511P, FL-B511M, FL-B512AG, FL-B512AP, FL-B512AM, FL-B513AG, FL-B513AP, FL-B513AM)

K260594 · Product code: **OHT** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k260594/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Based Over-the-counter Hair Removal (OHT)
Date received	Feb 23, 2026
Decision date	May 20, 2026
Days to decision	86 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Shenzhen Xiazhifeng Electronic Co., Ltd.**
Location	Shenzhen, CN
Contact	Meirong Li
510(k) history	2 submissions · 2 cleared · 2023-2026

REGULATORY CONSULTANT

Consulting firm	**Feiying Drug & Medical Consulting Technical Service Group**
Contact	Candice Qiu

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov