CLEARED

Oct 14, 2025
89 days to decision

K252234 IPL Home Use Hair Removal Device (Models: FDA11, FDA12, FDA13, FDA14, FDA15, FDA16, FDA17, FDA18, FDA19, FDA20, FDA21S, FDA22S, FDA23S, FDA24S, FDA25, FDA26, FDA27, FDA28, FDA29S, FDA30S, FDA31S)

K252234 · Product code: **OHT** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k252234/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Based Over-the-counter Hair Removal (OHT)
Date received	Jul 17, 2025
Decision date	Oct 14, 2025
Days to decision	89 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Shenzhen Qiaochengli Technology Co., Ltd.**
Location	Shenzhen, CN
Contact	Quanhua Huang
510(k) history	3 submissions · 3 cleared · 2021-2025

REGULATORY CONSULTANT

Consulting firm	**Feiying Drug & Medical Consulting Technical Service Group**
Contact	Riley Chen

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