

K230549 IPL Hair Removal Device, Model(s): FY-B505AL, FY-B505AG, FY-B505BL, FY-B505BG, FY-B507AL, FY-B507AG, FY-B507BL, FY-B507BG, FY-B508AL, FY-B508AG, FY-B508BL, FY-B508BG, FY-B509AL, FY-B509AG, FY-B509BL, FY-B509BGApr 25, 2023
56 days to decisionK230549 · Product code: OHT · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k230549/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Based Over-the-counter Hair Removal (OHT)
Date received	Feb 28, 2023
Decision date	Apr 25, 2023
Days to decision	56 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	James Li
510(k) history	1 submissions · 1 cleared · 2023-2023

REGULATORY CONSULTANT

Consulting firm	Feiyong Drug & Medical Consulting Technical Service Group
Contact	Tracy Che

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k230549/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026