FDA 510(k) Premarket Notification

CLEARED

K220222 IPL Hair Removal Device, model(s): S1-A, S2-A, S1, S2, S3, S4

Apr 26, 2022
90 days to decision

K220222 · Product code: **OHT** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k220222/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Based Over-the-counter Hair Removal (OHT)
Date received	Jan 26, 2022
Decision date	Apr 26, 2022
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Shenzhen Yuwei Electronic Technology Co., Ltd.**
Location	Shenzhen, CN
Contact	Liu Xuemeng
510(k) history	1 submissions · 1 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	**Feiying Drug & Medical Consulting Technical Service Group**
Contact	Rain Yip

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