FDA 510(k) Premarket Notification

CLEARED

K244020 Wrinkle Treatment Device (JM1, JM2B)

May 1, 2025
125 days to decision

K244020 · Product code: **OHS** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k244020/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Based Over The Counter Wrinkle Reduction (OHS)
Date received	Dec 27, 2024
Decision date	May 1, 2025
Days to decision	125 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Shenzhen Qianyu Technology Co., Ltd.**
Location	Shenzhen, CN
Contact	Guoyang Li
510(k) history	6 submissions · 6 cleared · 2022-2025

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k244020/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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