FDA 510(k) Premarket Notification

CLEARED

K260129 Lumaflex Panel (ZLD-05, ZLD-05A, ZLD-05APro)

Apr 15, 2026
89 days to decision

K260129 · Product code: **OLI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k260129/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fat Reducing Low Level Laser (OLI)
Date received	Jan 16, 2026
Decision date	Apr 15, 2026
Days to decision	89 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Shenzhen Kaiyan Medical Equipment Co., Ltd.**
Location	Shenzhen, CN
Contact	Alain Dijkstra
Website	https://www.kaiyanmedical.com
510(k) history	21 submissions · 21 cleared · 2023-2026

Shenzhen Kaiyan Medical Equipment Co., Ltd. is a medical device manufacturer based in Shenzhen, China. The company specializes in light therapy devices for clinical and aesthetic applications. Kaiyan Medical has received FDA 510(k) clearances from total submissions since 2023. All cleared devices fall within the General & Plastic Surgery category. The company maintains active regulatory status, with its most recent clearance in 2026. The company's cleared device portfolio includes LED light therapy masks, hair growth helmets, and advanced light-based aesthetic treatment s...