

K124043 LIGHTLAS FAMILY OF MULTI-WAVELENGTH MEDICAL LASER SYSTEM

Sep 26, 2013
272 days to decision

K124043 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k124043/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Dec 28, 2012
Decision date	Sep 26, 2013
Days to decision	272 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lightmed Corp.
Location	Shulin City, TW
Contact	JOCELYN LIU
510(k) history	9 submissions · 9 cleared · 2001-2015

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k124043/> 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 June 5, 2026