FDA 510(k) Premarket Notification

K124043 LIGHTLAS FAMILY OF MULTI-WAVELENGTH MEDICAL LASER SYSTEM

CLEARED

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