

**K091534 LIGHTLAS MODEL 532**Sep 18, 2009  
115 days to decisionK091534 · Product code: **HQF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k091534/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	May 26, 2009
Decision date	Sep 18, 2009
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Lightmed Corp.</b>
Location	Shulin City, TW
Contact	ANITA CHEN
510(k) history	9 submissions · 9 cleared · 2001-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091534/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 5, 2026