

**K181430 LENSAR Laser System - fs 3D (LLS-fs 3D)**Aug 9, 2018  
69 days to decisionK181430 · Product code: **OOE** · Ophthalmic  
Source: <https://www.510kdatabase.net/k181430/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ophthalmic Femtosecond Laser (OOE)
Date received	Jun 1, 2018
Decision date	Aug 9, 2018
Days to decision	69 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lensar, Inc.</b>
Location	Winter Park, FL, US
Contact	Keith Peck
510(k) history	14 submissions · 14 cleared · 2010-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k181430/> 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 May 27, 2026