

**K122829 LENSAR LASER SYSTEM - FS 3D FOR CORNEAL  
INCISIONS**Dec 3, 2012  
77 days to decisionK122829 · Product code: **HQC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k122829/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Sep 17, 2012
Decision date	Dec 3, 2012
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lensar, Inc.</b>
Location	Winter Park, FL, US
Contact	SHIRLEY K MCGARVEY
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/k122829/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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