

**K091915 FLUID ISOLATION DEVICE**Oct 27, 2009  
119 days to decisionK091915 · Product code: **HQC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k091915/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Jun 30, 2009
Decision date	Oct 27, 2009
Days to decision	119 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lenstec, Inc.</b>
Location	St. Petersburg, FL, US
Contact	JIMMY CHACKO
510(k) history	6 submissions · 6 cleared · 2005-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091915/> 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 2, 2026