

**K090938 FLOW CONTROLLER**Jun 4, 2009  
62 days to decisionK090938 · Product code: **LZU** · Ophthalmic  
Source: <https://www.510kdatabase.net/k090938/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plug, Punctum (LZU)
Date received	Apr 3, 2009
Decision date	Jun 4, 2009
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Odyssey Medical, Inc.</b>
Location	Memphis, TN, US
Contact	TERRY R GREEN
510(k) history	4 submissions · 4 cleared · 1997-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090938/> 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 23, 2026