

**K093510 LUMENVU CATHETER GUIDANCE SYSTEM**Aug 23, 2010  
284 days to decisionK093510 · Product code: **OMF** · General Hospital  
Source: <https://www.510kdatabase.net/k093510/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percutaneous, Implanted, Long-term Intravascular Catheter Accessory For Catheter Position (OMF)
Date received	Nov 12, 2009
Decision date	Aug 23, 2010
Days to decision	284 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sonosite, Inc.</b>
Location	Bothell, WA, US
Contact	JANICE HOGAN
510(k) history	23 submissions · 23 cleared · 1998-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093510/> 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 17, 2026