

**K871546 SAFESIDE(TM) LUMED**Jun 17, 1987  
58 days to decisionK871546 · Product code: **FOZ** · General Hospital  
Source: <https://www.510kdatabase.net/k871546/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Apr 20, 1987
Decision date	Jun 17, 1987
Days to decision	58 days
Third-party review	No

**APPLICANT**

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Company	<b>Luther Medical Products, Inc.</b>
Location	Walker, MI, US
Contact	RONALD B LUTHER
510(k) history	17 submissions · 16 cleared · 1980-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k871546/>; 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 July 3, 2026