

**K883768 VAS-CATH CATHETER REPAIR KIT**Nov 14, 1988  
69 days to decisionK883768 · Product code: **LFJ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k883768/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Sep 6, 1988
Decision date	Nov 14, 1988
Days to decision	69 days
Third-party review	No

**APPLICANT**

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Company	<b>Vas-Cath, Inc.</b>
Location	Mississauga, Ontario, CA
Contact	H CREWE
510(k) history	27 submissions · 27 cleared · 1988-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883768/>; 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 1, 2026