

**K875188 LUER LOCK CONNECTORS / RIDIG P.V.C.**Mar 2, 1988  
76 days to decisionK875188 · Product code: **LFJ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k875188/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Dec 17, 1987
Decision date	Mar 2, 1988
Days to decision	76 days
Third-party review	No

**APPLICANT**

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Company	<b>Vas-Cath of Canada , Ltd.</b>
Location	Mchenry, IL, US
Contact	KATHERINE CREWE
510(k) history	19 submissions · 19 cleared · 1979-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k875188/>; 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 27, 2026