

**K892926 DUAL LUMEN CATHETER FOR CHRONIC VASCULAR ACCESS**Dec 5, 1989  
228 days to decisionK892926 · Product code: **LJS** · General Hospital  
Source: <https://www.510kdatabase.net/k892926/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Apr 21, 1989
Decision date	Dec 5, 1989
Days to decision	228 days
Third-party review	No

**APPLICANT**

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Company	<b>Vas-Cath, Inc.</b>
Location	Mississauga, Ontario, CA
Contact	KATHERINE 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/k892926/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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