

**K895506 FLEXXICON PLUS DUAL LUMEN DIALYSIS  
CATHETER**Feb 9, 1990  
150 days to decisionK895506 · Product code: **LFJ** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k895506/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Sep 12, 1989
Decision date	Feb 9, 1990
Days to decision	150 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/k895506/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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