

**K891753 DUAL LUMEN SUB-CLAVIAN CATHETER, DLSC 600 & 800**Jun 8, 1989  
77 days to decisionK891753 · Product code: **LFJ** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k891753/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Mar 23, 1989
Decision date	Jun 8, 1989
Days to decision	77 days
Third-party review	No

**APPLICANT**

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Company	<b>Akcess Medical Products, Inc.</b>
Location	New Brunswick, NJ, US
Contact	EDWARD J HOPKINS
510(k) history	36 submissions · 28 cleared · 1989-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k891753/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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