

**K881695 DESERET SUBCLAVIAN HEMODIALYSIS CATHETER**Jun 23, 1988  
65 days to decisionK881695 · Product code: **LFJ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k881695/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Apr 19, 1988
Decision date	Jun 23, 1988
Days to decision	65 days
Third-party review	No

**APPLICANT**

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Company	<b>Deseret Medical, Inc.</b>
Location	Mchenry, IL, US
Contact	CHARLES J WELLE
510(k) history	20 submissions · 20 cleared · 1979-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881695/>; 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