

K833848 DUAL LUMEN SUBCLAVIAN CANNULADec 8, 1983
34 days to decisionK833848 · Product code: **LFJ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k833848/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Nov 4, 1983
Decision date	Dec 8, 1983
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Shiley, Inc.
Location	Mchenry, IL, US
510(k) history	174 submissions · 174 cleared · 1976-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k833848/> Data sourced from FDA 510(k) public records (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. - Generated May 26, 2026