

K881290 MODIFIED SHILEY DUAL LUMEN SUBCLAVIAN CANNULAApr 27, 1988
30 days to decisionK881290 · Product code: **LFJ** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k881290/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Mar 28, 1988
Decision date	Apr 27, 1988
Days to decision	30 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881290/> 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