

**K895417 ARROW LARGE-BORE DUAL LUMEN HEMODIALYSIS
KIT**Jan 3, 1990
124 days to decisionK895417 · Product code: **LFJ** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k895417/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Sep 1, 1989
Decision date	Jan 3, 1990
Days to decision	124 days
Third-party review	No

APPLICANT

Company	Arrow Intl., Inc.
Location	Mchenry, IL, US
Contact	THOMAS D NICKEL
510(k) history	110 submissions · 105 cleared · 1976-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895417/>; 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 22, 2026