FDA 510(k) Premarket Notification

K895417 ARROW LARGE-BORE DUAL LUMEN HEMODIALYSIS KIT

CLEARED

Jan 3, 1990
124 days to decision

K895417 · 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