FDA 510(k) Premarket Notification

K900263 ANTIMICROBIAL MULTI-LUMEN CENTRAL VENOUS CATHETER

CLEARED

Jul 24, 1990
187 days to decision

K900263 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k900263/

SUBMISSION DETAILS

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Jan 18, 1990
Decision date	Jul 24, 1990
Days to decision	187 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