FDA 510(k) Premarket Notification

K862056 [CENTRAL VENOUS][MULTI-LUMEN] CATHETER W/FLEX TIP

CLEARED

Aug 25, 1986
87 days to decision

K862056 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k862056/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	May 30, 1986
Decision date	Aug 25, 1986
Days to decision	87 days
Third-party review	No

APPLICANT

Company	**Arrow Intl., Inc.**
Location	Mchenry, IL, US
Contact	RONALD P CITRON
510(k) history	110 submissions · 105 cleared · 1976-2010