FDA 510(k) Premarket Notification

K960479 ARROW BIPOLAR PACING/BALLOON WEDGE PRESSURE CATHETER

CLEARED

Oct 17, 1996
258 days to decision

K960479 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k960479/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Feb 2, 1996
Decision date	Oct 17, 1996
Days to decision	258 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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