FDA 510(k) Premarket Notification

K040100 ARROW ECHOGENIC INTRODUCER NEEDLE

CLEARED

Mar 1, 2004
41 days to decision

K040100 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k040100/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jan 20, 2004
Decision date	Mar 1, 2004
Days to decision	41 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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