FDA 510(k) Premarket Notification

K901899 ROADRUNNER WIRE GUIDE

CLEARED

Sep 27, 1990
153 days to decision

K901899 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k901899/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Apr 27, 1990
Decision date	Sep 27, 1990
Days to decision	153 days
Third-party review	No

APPLICANT

Company	**Cook, Inc.**
Location	Mchenry, IL, US
Contact	APRIL LAVENDER
510(k) history	190 submissions · 179 cleared · 1976-2015