FDA 510(k) Premarket Notification

K920891 ROADRUNNER(TM) WIRE GUIDE, MODIFICATION

CLEARED

Dec 21, 1992
301 days to decision

K920891 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k920891/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 24, 1992
Decision date	Dec 21, 1992
Days to decision	301 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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