FDA 510(k) Premarket Notification

K121776 AGILITY STEERABLE GUIDEWIRE NEUROSCOUT STEERABLE GUDIEWIRE

CLEARED

Aug 14, 2012
57 days to decision

K121776 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k121776/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 18, 2012
Decision date	Aug 14, 2012
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Codman & Shurtleff, Inc.**
Location	Mchenry, IL, US
Contact	JOAN BARTLE
510(k) history	152 submissions · 151 cleared · 1976-2020