FDA 510(k) Premarket Notification

K100351 NEUROSCOUT STEERABLE GUIDEWIRE

CLEARED

Mar 16, 2010
32 days to decision

K100351 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k100351/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 12, 2010
Decision date	Mar 16, 2010
Days to decision	32 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